

# Prior Authorization Criteria

## ABATACEPT IV

### Products Affected

- ORENCIA INTRAVENOUS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA), POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST.
Coverage Duration	RA, PJIA, PSA: INITIAL: 6 MOS, RENEWAL: 12 MOS. ACUTE GRAFT VERSUS HOST DISEASE (AGVHD): 1 MO.
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. PJIA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PJIA. PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. RENEWAL: RA: CONTINUES TO BENEFIT FROM THE MEDICATION. PJIA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PJIA. PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA.

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ABATACEPT SQ

## Products Affected

- ORENCIA CLICKJECT
- ORENCIA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA), POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. PJIA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PJIA. PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. RENEWAL: RA: CONTINUES TO BENEFIT FROM THE MEDICATION. PJIA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PJIA. PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA.

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# ABEMACICLIB

---

## Products Affected

- VERZENIO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ABIRATERONE

## Products Affected

- *abiraterone acetate*
- *abirtega*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC HIGH-RISK CASTRATION-SENSITIVE PROSTATE CANCER (MCSPC), METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC): 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# ABIRATERONE SUBMICRONIZED

**Products Affected**

- YONSA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC): 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# ACALABRUTINIB

## Products Affected

- CALQUENCE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PREVIOUSLY TREATED MANTLE CELL LYMPHOMA: INTOLERANCE TO BRUKINSA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ADAGRASIB

---

## Products Affected

- KRAZATI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ADALIMUMAB

## Products Affected

- HUMIRA (2 PEN) SUBCUTANEOUS AUTO-INJECTOR KIT
- HUMIRA (2 SYRINGE) SUBCUTANEOUS PREFILLED SYRINGE KIT 10 MG/0.1ML, 20 MG/0.2ML, 40 MG/0.4ML, 40 MG/0.8ML
- HUMIRA-CD/UC/HS STARTER SUBCUTANEOUS AUTO-INJECTOR KIT
- HUMIRA-PED<40KG CROHNS STARTER
- HUMIRA-PED>=40KG CROHNS START
- HUMIRA-PED>=40KG UC STARTER SUBCUTANEOUS AUTO-INJECTOR KIT
- HUMIRA-PS/UV/ADOL HS STARTER SUBCUTANEOUS AUTO-INJECTOR KIT
- HUMIRA-PSORIASIS/UVEIT STARTER SUBCUTANEOUS AUTO-INJECTOR KIT

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP, OR GENITAL AREA.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: RA, PJIA, ANKYLOSING SPONDYLITIS (AS): PRESCRIBED BY OR IN CONSULTATION WITH RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH DERMATOLOGIST OR RHEUMATOLOGIST. PSO, HIDRADENITIS SUPPURATIVA (HS): PRESCRIBED BY OR IN CONSULTATION WITH DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH GASTROENTEROLOGIST. UVEITIS: PRESCRIBED BY OR IN CONSULTATION WITH OPHTHALMOLOGIST
<b>Coverage Duration</b>	INITIAL: RA, PSO, PJIA, AS, PSA, CD, UC, UVEITIS: 6 MONTHS, HS: 12 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: RHEUMATOID ARTHRITIS (RA): TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED.</p> <p>POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PJIA. PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. AS: 1) TRIAL OF OR CONTRAINDICATION TO AN NSAID AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. PSO: 1) ONE OF THE FOLLOWING: (A) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY) FOR THE TREATMENT OF PSO, (B) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA FOR THE TREATMENT OF PSO, OR (C) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC. HS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR HS. UVEITIS: NO ISOLATED ANTERIOR UVEITIS. RENEWAL: RA, HS, UVEITIS: CONTINUES TO BENEFIT FROM THE MEDICATION. PJIA: 1) CONTINUES TO BENEFIT FROM MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PJIA. PSA: 1) CONTINUES TO BENEFIT FROM MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. AS: 1) CONTINUES TO BENEFIT FROM MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. PSO: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO</p>

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
	CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ADALIMUMAB-AATY

## Products Affected

- YUFLYMA (1 PEN)
- YUFLYMA (2 SYRINGE)
- YUFLYMA-CD/UC/HS STARTER

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP, OR GENITAL AREA.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: RA, PJA, ANKYLOSING SPONDYLITIS (AS): PRESCRIBED BY OR IN CONSULTATION WITH RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH DERMATOLOGIST OR RHEUMATOLOGIST. PSO, HIDRADENITIS SUPPURATIVA (HS): PRESCRIBED BY OR IN CONSULTATION WITH DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH GASTROENTEROLOGIST. UVEITIS: PRESCRIBED BY OR IN CONSULTATION WITH OPHTHALMOLOGIST
<b>Coverage Duration</b>	INITIAL: RA, PSO, PJA, AS, PSA, CD, UC, UVEITIS: 6 MONTHS, HS: 12 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: RHEUMATOID ARTHRITIS (RA): TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJA): NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PJA. PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. AS: 1) TRIAL OF OR

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>CONTRAINDICATION TO AN NSAID AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. PSO: 1) ONE OF THE FOLLOWING: (A) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY) FOR THE TREATMENT OF PSO, (B) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA FOR THE TREATMENT OF PSO, OR (C) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC. HS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR HS. UVEITIS: NO ISOLATED ANTERIOR UVEITIS. RENEWAL: RA, HS, UVEITIS: CONTINUES TO BENEFIT FROM MEDICATION. PJIA: 1) CONTINUES TO BENEFIT FROM MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PJIA. PSA: 1) CONTINUES TO BENEFIT FROM MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. AS: 1) CONTINUES TO BENEFIT FROM MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. PSO: 1) CONTINUES TO BENEFIT FROM MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC.</p>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ADALIMUMAB-ADBIM

## Products Affected

- CYLTEZO (2 PEN)
- CYLTEZO (2 SYRINGE)
- CYLTEZO-CD/UC/HS STARTER
- CYLTEZO-PSORIASIS/UV STARTER

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP, OR GENITAL AREA.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: RA, PJIA, ANKYLOSING SPONDYLITIS (AS): PRESCRIBED BY OR IN CONSULTATION WITH RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH DERMATOLOGIST OR RHEUMATOLOGIST. PSO, HIDRADENITIS SUPPURATIVA (HS): PRESCRIBED BY OR IN CONSULTATION WITH DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH GASTROENTEROLOGIST. UVEITIS: PRESCRIBED BY OR IN CONSULTATION WITH OPHTHALMOLOGIST
<b>Coverage Duration</b>	INITIAL: RA, PSO, PJIA, AS, PSA, CD, UC, UVEITIS: 6 MONTHS, HS: 12 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: RHEUMATOID ARTHRITIS (RA): TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PJIA. PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>MOLECULES FOR PSA. AS: 1) TRIAL OF OR CONTRAINDICATION TO AN NSAID AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. PSO: 1) ONE OF THE FOLLOWING: (A) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY) FOR THE TREATMENT OF PSO, (B) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA FOR THE TREATMENT OF PSO, OR (C) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC. HS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR HS. UVEITIS: NO ISOLATED ANTERIOR UVEITIS. RENEWAL: RA, HS, UVEITIS: CONTINUES TO BENEFIT FROM MEDICATION. PJIA: 1) CONTINUES TO BENEFIT FROM MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PJIA. PSA: 1) CONTINUES TO BENEFIT FROM MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. AS: 1) CONTINUES TO BENEFIT FROM MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. PSO: 1) CONTINUES TO BENEFIT FROM MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC.</p>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# AFATINIB

---

## Products Affected

- GILOTRIF

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EGFR MUTATION: NOT ON CONCURRENT THERAPY WITH AN EGFR TYROSINE KINASE INHIBITOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ALECTINIB

---

## Products Affected

- ALECENSA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ALPELISIB-PIQRAY

## Products Affected

- PIQRAY (200 MG DAILY DOSE)
- PIQRAY (250 MG DAILY DOSE)
- PIQRAY (300 MG DAILY DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# AMIKACIN LIPOSOMAL INH

## Products Affected

- ARIKAYCE

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	MYCOBACTERIUM AVIUM COMPLEX (MAC) LUNG DISEASE: RENEWAL: 1) NO POSITIVE MAC SPUTUM CULTURE AFTER CONSECUTIVE NEGATIVE CULTURES, AND 2) IMPROVEMENT IN SYMPTOMS. ADDITIONALLY, FOR FIRST RENEWAL, APPROVAL REQUIRES AT LEAST ONE NEGATIVE SPUTUM CULTURE FOR MAC BY SIX MONTHS OF ARIKAYCE TREATMENT. FOR SECOND AND SUBSEQUENT RENEWALS, APPROVAL REQUIRES AT LEAST THREE NEGATIVE SPUTUM CULTURES FOR MAC BY 12 MONTHS OF ARIKAYCE TREATMENT.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	MAC LUNG DISEASE: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR INFECTIOUS DISEASE SPECIALIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 6 MONTHS.
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# AMIVANTAMAB-VMJW

## Products Affected

- RYBREVANT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# ANAKINRA

## Products Affected

- KINERET SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	CORONAVIRUS DISEASE 2019 (COVID-19) IN HOSPITALIZED ADULTS.
<b>Required Medical Information</b>	INITIAL: CRYOPYRIN-ASSOCIATED PERIODIC SYNDROMES (CAPS): 1) ONE OF THE FOLLOWING: (A) GENETIC TEST FOR GAIN-OF-FUNCTION MUTATIONS IN THE NLRP3 GENE, OR (B) HAS INFLAMMATORY MARKERS (I.E., ELEVATED CRP, ESR, SERUM AMYLOID A PROTEIN (SAA) OR S100 PROTEINS), AND 2) TWO OF THE FOLLOWING: URTICARIAL-LIKE RASH (NEUTROPHILIC DERMATITIS), COLD-TRIGGERED EPISODES, SENSORINEURAL HEARING LOSS, MUSCULOSKELETAL SYMPTOMS, CHRONIC ASEPTIC MENINGITIS, SKELETAL ABNORMALITIES. DEFICIENCY OF INTERLEUKIN-1 RECEPTOR ANTAGONIST (DIRA): 1) ONE OF THE FOLLOWING: (A) GENETIC TEST FOR GAIN-OF-FUNCTION MUTATIONS IN THE IL1RN GENE, OR (B) HAS INFLAMMATORY MARKERS (I.E., ELEVATED CRP, ESR), AND 2) ONE OF THE FOLLOWING: PUSTULAR PSORIASIS-LIKE RASHES, OSTEOMYELITIS, ABSENCE OF BACTERIAL OSTEOMYELITIS, ONYCHOMADESIS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: RHEUMATOID ARTHRITIS (RA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST.
<b>Coverage Duration</b>	RA: INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS. CAPS, DIRA: LIFETIME.
<b>Other Criteria</b>	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ, RINVOQ, ORENCIA. CAPS, DIRA: NO CONCURRENT USE WITH OTHER IL-1 INHIBITORS. RENEWAL: RA: CONTINUES TO BENEFIT FROM THE MEDICATION.
<b>Indications</b>	All FDA-approved Indications.

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# APALUTAMIDE

## Products Affected

- ERLEADA ORAL TABLET 240 MG, 60 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: NON-METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (NMCRPC): HIGH RISK PROSTATE CANCER (I.E., RAPIDLY INCREASING PROSTATE SPECIFIC ANTIGEN [PSA] LEVELS). NMCRPC, METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (MCSPC): 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG. RENEWAL: NMCRPC, MCSPC: 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GNRH ANALOG.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# APOMORPHINE - ONAPGO

## Products Affected

- ONAPGO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PARKINSONS DISEASE (PD): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL: 3 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	PD: INITIAL: 1) RESPONSIVE TO LEVODOPA, 2) CURRENT MEDICATION REGIMEN, INCLUDING LEVODOPA, HAS BEEN AT A STABLE DOSE FOR AT LEAST 28 DAYS, 3) MOTOR SYMPTOMS ARE CURRENTLY UNCONTROLLED (DEFINED AS AN AVERAGE OFF TIME OF AT LEAST 3 HOURS/DAY, FOR AT LEAST 2 HOURS EACH DAY), AND 4) DOES NOT HAVE ANY OF THE FOLLOWING: ORTHOSTATIC HYPOTENSION, HISTORY OF PROLONGED QTC (GREATER THAN 450 MSEC FOR MALE OR GREATER THAN 470 MSEC FOR FEMALE), ACTIVE OR UNCONTROLLED PSYCHOSIS, ACTIVE OR UNCONTROLLED DEPRESSION. RENEWAL: IMPROVEMENT IN MOTOR SYMPTOMS WHILE ON THERAPY. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# APOMORPHINE - SL

## Products Affected

- KYNMOBI
- KYNMOBI TITRATION KIT

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	PARKINSONS DISEASE (PD): INITIAL: 18 YEARS OF AGE OR OLDER.
<b>Prescriber Restrictions</b>	PD: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
<b>Other Criteria</b>	PD: RENEWAL: IMPROVEMENT WITH MOTOR FLUCTUATIONS DURING OFF EPISODES WITH THE USE OF THERAPY.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# APREMILAST

## Products Affected

- OTEZLA

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: MILD PLAQUE PSORIASIS (PSO): 1) PSORIASIS COVERING 2 PERCENT OF BODY SURFACE AREA (BSA), 2) STATIC PHYSICIAN GLOBAL ASSESSMENT (SPGA) SCORE OF 2, OR 3) PSORIASIS AREA AND SEVERITY INDEX (PASI) SCORE OF 2 TO 9. MODERATE TO SEVERE PSO: PSORIASIS COVERING 3 PERCENT OR MORE OF BSA, OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP, OR GENITAL AREA.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. BEHCETS DISEASE: PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PSA. MILD PSO: TRIAL OF OR CONTRAINDICATION TO ONE CONVENTIONAL SYSTEMIC THERAPY (E.G., METHOTREXATE, ACITRETIN, CYCLOSPORINE) OR ONE CONVENTIONAL TOPICAL THERAPY (E.G., PUVA [PHOTOTHERAPY], UVB [ULTRAVIOLET LIGHT B], TOPICAL CORTICOSTEROIDS). MODERATE TO SEVERE PSO: 1) ONE OF THE FOLLOWING: (A) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY) FOR THE TREATMENT OF PSO, (B) CONTRAINDICATION OR

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA FOR THE TREATMENT OF PSO, OR (C) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR MODERATE TO SEVERE PSO. BEHCETS DISEASE: 1) HAS ORAL ULCERS OR A HISTORY OF RECURRENT ORAL ULCERS BASED ON CLINICAL SYMPTOMS, AND 2) TRIAL OF OR CONTRAINDICATION TO ONE OR MORE CONSERVATIVE TREATMENTS (E.G., COLCHICINE, TOPICAL CORTICOSTEROID, ORAL CORTICOSTEROID). RENEWAL: MILD PSO, BEHCETS DISEASE: CONTINUES TO BENEFIT FROM THE MEDICATION. PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. MODERATE TO SEVERE PSO: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR MODERATE TO SEVERE PSO.</p>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ARIMOCLOMOL

## Products Affected

- MIPLYFFA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	NIEMANN-PICK DISEASE TYPE C (NPC): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH NEUROLOGIST OR GENETICIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	NPC: RENEWAL: IMPROVEMENT OR SLOWING OF DISEASE PROGRESSION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ASCIMINIB

## Products Affected

- SCEMBLIX ORAL TABLET 100 MG, 20 MG, 40 MG

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	PREVIOUSLY TREATED OR T315I MUTATION PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (Ph+ CML): MUTATIONAL ANALYSIS PRIOR TO INITIATION AND SCEMBLIX IS APPROPRIATE PER NCCN GUIDELINE TABLE FOR TREATMENT RECOMMENDATIONS BASED ON BCR-ABL1 MUTATION PROFILE.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# ASFOTASE ALFA

## Products Affected

- STRENSIQ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	HYPOPHOSPHATASIA (HPP): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST, GENETICIST, OR METABOLIC SPECIALIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: PERINATAL/INFANTILE-ONSET HPP: 1) 6 MONTHS OF AGE OR YOUNGER AT ONSET OF HPP, AND 2) POSITIVE FOR A TISSUE NON-SPECIFIC ALKALINE PHOSPHATASE (TNSALP) (ALPL) GENE MUTATION AS CONFIRMED BY GENETIC TESTING OR TWO OF THE FOLLOWING: (A) SERUM ALKALINE PHOSPHATASE (ALP) LEVEL BELOW THAT OF NORMAL RANGE FOR PATIENT AGE, (B) ELEVATED SERUM PYRIDOXAL-5'-PHOSPHATE (PLP) LEVELS AND NO VITAMIN B6 SUPPLEMENTATION IN THE PREVIOUS WEEK, (C) URINE PHOSPHOETHANOLAMINE (PEA) LEVEL ABOVE THAT OF NORMAL RANGE FOR PATIENT AGE, (D) RADIOGRAPHIC EVIDENCE OF HPP, (E) AT LEAST TWO OF THE FOLLOWING: (I) RACHITIC CHEST DEFORMITY, (II) CRANIOSYNOSTOSIS, (III) DELAY IN SKELETAL GROWTH RESULTING IN DELAY OF MOTOR DEVELOPMENT, (IV) HISTORY OF VITAMIN B6 DEPENDENT SEIZURES, (V) NEPHROCALCINOSIS OR HISTORY OF ELEVATED SERUM CALCIUM, (VI) HISTORY OR PRESENCE OF NON-TRAUMATIC POSTNATAL FRACTURE AND DELAYED FRACTURE HEALING. JUVENILE-ONSET HPP: 1) 18 YEARS OF AGE OR YOUNGER AT ONSET OF HPP, AND 2) POSITIVE FOR A

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>TNSALP ALPL GENE MUTATION AS CONFIRMED BY GENETIC TESTING OR TWO OF THE FOLLOWING: (A) SERUM ALP LEVEL BELOW THAT OF NORMAL RANGE FOR PATIENT AGE, (B) ELEVATED SERUM PLP LEVELS AND NO VITAMIN B6 SUPPLEMENTATION IN THE PREVIOUS WEEK, (C) URINE PEA LEVEL ABOVE THAT OF NORMAL RANGE FOR PATIENT AGE, (D) RADIOGRAPHIC EVIDENCE OF HPP, (E) AT LEAST TWO OF THE FOLLOWING: (I) RACHITIC DEFORMITIES, (II) PREMATURE LOSS OF PRIMARY TEETH PRIOR TO 5 YEARS OF AGE, (III) DELAY IN SKELETAL GROWTH RESULTING IN DELAY OF MOTOR DEVELOPMENT, (IV) HISTORY OR PRESENCE OF NON-TRAUMATIC FRACTURES OR DELAYED FRACTURE HEALING. ALL INDICATIONS: 1) NOT CURRENTLY RECEIVING TREATMENT WITH A BISPHOSPHONATE, 2) CALCIUM OR PHOSPHATE LEVELS ARE NOT BELOW THE NORMAL RANGE, 3) NOT HAVE A TREATABLE FORM OF RICKETS. RENEWAL: ALL INDICATIONS: 1) IMPROVEMENT IN THE SKELETAL CHARACTERISTICS OF HPP, AND 2) NOT CURRENTLY RECEIVING TREATMENT WITH A BISPHOSPHONATE.</p>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ATOGEPANT

## Products Affected

- QULIPTA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	MIGRAINE PREVENTION: INITIAL: 1) TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING PREVENTIVE MIGRAINE TREATMENTS: DIVALPROEX SODIUM, TOPIRAMATE, PROPRANOLOL, TIMOLOL, AND 2) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR MIGRAINE PREVENTION. RENEWAL: 1) REDUCTION IN MIGRAINE OR HEADACHE FREQUENCY, MIGRAINE SEVERITY, OR MIGRAINE DURATION WITH THERAPY, AND 2) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR MIGRAINE PREVENTION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# AVACOPAN

## Products Affected

- TAVNEOS

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	ANTI-NEUTROPHIL CYTOPLASMIC AUTOANTIBODY (ANCA)-ASSOCIATED VASCULITIS: INITIAL: ANCA SEROPOSITIVE (ANTI-PR3 OR ANTI-MPO).
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	ANCA-ASSOCIATED VASCULITIS: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR NEPHROLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 6 MONTHS.
<b>Other Criteria</b>	ANCA-ASSOCIATED VASCULITIS: RENEWAL: CONTINUES TO BENEFIT FROM THERAPY.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# AVAPRITINIB

---

## Products Affected

- AYVAKIT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# AXATILIMAB-CSFR

---

## Products Affected

- NIKTIMVO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# AXITINIB

---

## Products Affected

- INLYTA ORAL TABLET 1 MG, 5 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# AZACITIDINE

---

## Products Affected

- ONUREG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# AZTREONAM INHALED

---

## Products Affected

- CAYSTON

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	7 YEARS OF AGE OR OLDER
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# BEDAQUILINE

## Products Affected

- SIRTURO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	24 WEEKS
Other Criteria	PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS (MDR-TB); SIRTURO USED IN COMBINATION WITH AT LEAST 3 OTHER ANTIBIOTICS FOR THE TREATMENT OF PULMONARY MDR-TB.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# BELIMUMAB

## Products Affected

- BENLYSTA SUBCUTANEOUS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: SYSTEMIC LUPUS ERYTHEMATOSUS (SLE): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. LUPUS NEPHRITIS (LN): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR NEPHROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: SLE: CURRENTLY TAKING CORTICOSTEROIDS, ANTIMALARIALS, NSAIDS, OR IMMUNOSUPPRESSIVE AGENTS. RENEWAL: SLE: PATIENT HAD CLINICAL IMPROVEMENT. LN: IMPROVEMENT IN RENAL RESPONSE FROM BASELINE LABORATORY VALUES (I.E., EGFR OR PROTEINURIA) AND/OR CLINICAL PARAMETERS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# BELUMOSUDIL

---

## Products Affected

- REZUROCK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# BELZUTIFAN

---

## Products Affected

- WELIREG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# BENDAMUSTINE

**Products Affected**

- BENDAMUSTINE HCL INTRAVENOUS SOLUTION
- *bendamustine hcl intravenous solution reconstituted*
- BENDEKA
- VIVIMUSTA

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# BENRALIZUMAB

## Products Affected

- FASENRA
- FASENRA PEN

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	ASTHMA: INITIAL: BLOOD EOSINOPHIL LEVEL OF AT LEAST 150 CELLS/MCL WITHIN THE PAST 12 MONTHS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	ASTHMA: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A PHYSICIAN SPECIALIZING IN ALLERGY OR PULMONARY MEDICINE.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: ASTHMA: 1) CONCURRENT THERAPY WITH A MEDIUM, HIGH-DOSE, OR MAXIMALLY TOLERATED DOSE OF AN INHALED CORTICOSTEROID (ICS) AND ONE OTHER MAINTENANCE MEDICATION, 2) ONE ASTHMA EXACERBATION REQUIRING SYSTEMIC CORTICOSTEROID BURST LASTING 3 OR MORE DAYS WITHIN THE PAST 12 MONTHS, OR AT LEAST ONE SERIOUS EXACERBATION REQUIRING HOSPITALIZATION OR ER VISIT WITHIN THE PAST 12 MONTHS, OR POOR SYMPTOM CONTROL DESPITE CURRENT THERAPY AS EVIDENCED BY AT LEAST THREE OF THE FOLLOWING WITHIN THE PAST 4 WEEKS: (A) DAYTIME ASTHMA SYMPTOMS MORE THAN TWICE/WEEK, (B) ANY NIGHT WAKING DUE TO ASTHMA, (C) SABA RELIEVER FOR SYMPTOMS MORE THAN TWICE/WEEK, (D) ANY ACTIVITY LIMITATION DUE TO ASTHMA, AND 3) NO CONCURRENT USE WITH XOLAIR, DUPIXENT, TEZSPIRE, OR OTHER ANTI-IL5 BIOLOGICS WHEN USED FOR ASTHMA. EOSINOPHILIC GRANULOMATOSIS WITH POLYANGIITIS (EGPA): NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-2

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	INHIBITOR) FOR EGPA. RENEWAL: ASTHMA: 1) NO CONCURRENT USE WITH XOLAIR, DUPIXENT, TEZSPIRE, OR OTHER ANTI-IL5 BIOLOGICS WHEN USED FOR ASTHMA, 2) CONTINUED USE OF ICS AND ONE OTHER MAINTENANCE MEDICATION, AND 3) CLINICAL RESPONSE AS EVIDENCED BY: (A) REDUCTION IN ASTHMA EXACERBATIONS FROM BASELINE, (B) DECREASED UTILIZATION OF RESCUE MEDICATIONS, (C) INCREASE IN PERCENT PREDICTED FEV1 FROM PRETREATMENT BASELINE, OR (D) REDUCTION IN SEVERITY OR FREQUENCY OF ASTHMA-RELATED SYMPTOMS. EGPA: 1) REDUCTION IN EGPA SYMPTOMS COMPARED TO BASELINE OR ABILITY TO REDUCE/ELIMINATE CORTICOSTEROID USE, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR EGPA.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# BETAINE

---

## Products Affected

- *betaine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# BEVACIZUMAB-ADCD

## Products Affected

- VEGZELMA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# BEVACIZUMAB-AWWB

## Products Affected

- MVASI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# BEVACIZUMAB-BVZR

## Products Affected

- ZIRABEV

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# BEXAROTENE

---

## Products Affected

- *bexarotene*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# BINIMETINIB

---

## Products Affected

- MEKTOVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# BORTEZOMIB

## Products Affected

- *bortezomib injection*
- BORUZU

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# BOSENTAN

## Products Affected

- *bosentan*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	PULMONARY ARTERIAL HYPERTENSION (PAH): INITIAL: DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	PAH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	PAH: INITIAL: 1) DOES NOT HAVE ELEVATED LIVER ENZYMES (ALT, AST) MORE THAN 3 TIMES UPPER LIMIT OF NORMAL (ULN) OR INCREASE IN BILIRUBIN BY 2 OR MORE TIMES ULN, AND 2) NO CONCURRENT USE WITH CYCLOSPORINE A OR GLYBURIDE. RENEWAL: NO CONCURRENT USE WITH CYCLOSPORINE A OR GLYBURIDE.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# BOSUTINIB

## Products Affected

- BOSULIF ORAL CAPSULE 100 MG, 50 MG
- BOSULIF ORAL TABLET 100 MG, 400 MG, 500 MG

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	PREVIOUSLY TREATED PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (Ph+ CML): MUTATIONAL ANALYSIS PRIOR TO INITIATION AND BOSULIF IS APPROPRIATE PER NCCN GUIDELINE TABLE FOR TREATMENT RECOMMENDATIONS BASED ON BCR-ABL1 MUTATION PROFILE.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# BRIGATINIB

## Products Affected

- ALUNBRIG ORAL TABLET 180 MG, 30 MG, 90 MG
- ALUNBRIG ORAL TABLET THERAPY PACK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# C1 ESTERASE INHIBITOR-HAEGARDA

## Products Affected

- HAEGARDA SUBCUTANEOUS SOLUTION RECONSTITUTED 2000 UNIT, 3000 UNIT

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	HEREDITARY ANGIOEDEMA (HAE): INITIAL: DIAGNOSIS CONFIRMED BY ONE OF THE FOLLOWING COMPLEMENT TESTING: C1INH PROTEIN LEVELS, C4 PROTEIN LEVELS, C1-INH FUNCTIONAL LEVELS, C1Q.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	HAE: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST, IMMUNOLOGIST, ALLERGIST OR PULMONOLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	HAE: INITIAL: NOT ON CONCURRENT TREATMENT WITH ALTERNATIVE PROPHYLACTIC AGENT FOR HAE ATTACKS. RENEWAL: 1) IMPROVEMENT COMPARED TO BASELINE IN HAE ATTACKS (I.E., REDUCTIONS IN ATTACK FREQUENCY OR ATTACK SEVERITY), AND 2) NOT ON CONCURRENT TREATMENT WITH ALTERNATIVE PROPHYLACTIC AGENT FOR HAE ATTACKS.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# CABOZANTINIB CAPSULE

## Products Affected

- COMETRIQ (100 MG DAILY DOSE)  
ORAL KIT 80 & 20 MG
- COMETRIQ (140 MG DAILY DOSE)  
ORAL KIT 3 X 20 MG & 80 MG
- COMETRIQ (60 MG DAILY DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# CABOZANTINIB TABLET

## Products Affected

- CABOMETYX ORAL TABLET 20 MG, 40 MG, 60 MG

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# CANNABIDIOL

## Products Affected

- EPIDIOLEX

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	DRAVET SYNDROME (DS), LENNOX-GASTAUT SYNDROME (LGS), TUBEROUS SCLEROSIS COMPLEX (TSC): PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: LENNOX-GASTAUT SYNDROME (LGS): TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING ANTIPILEPTIC MEDICATIONS: RUFINAMIDE, FELBAMATE, CLOBAZAM, TOPIRAMATE, LAMOTRIGINE, CLONAZEPAM.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# CAPIVASERTIB

## Products Affected

- TRUQAP ORAL TABLET
- TRUQAP TABLET THERAPY PACK  
160 MG ORAL

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# CAPMATINIB

---

## Products Affected

- TABRECTA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# CARGLUMIC ACID

## Products Affected

- carglumic acid oral tablet soluble*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: ACUTE OR CHRONIC HYPERAMMONEMIA (HA) DUE TO N ACETYLGLUTAMATE SYNTHASE (NAGS) DEFICIENCY: NAGS GENE MUTATION IS CONFIRMED BY BIOCHEMICAL OR GENETIC TESTING. ACUTE HA DUE TO PROPIONIC ACIDEMIA (PA): 1) CONFIRMED BY ELEVATED METHYLCITRIC ACID AND NORMAL METHYLMALONIC ACID, OR 2) GENETIC TESTING CONFIRMS MUTATION IN THE PCCA OR PCCB GENE. ACUTE HA DUE TO METHYLMALONIC ACIDEMIA (MMA): 1) CONFIRMED BY ELEVATED METHYLMALONIC ACID, METHYLCITRIC ACID, OR 2) GENETIC TESTING CONFIRMS MUTATION IN THE MMUT, MMA, MMAB OR MMADHC GENES.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	ACUTE HA DUE TO NAGS/PA/MMA: 7 DAYS. CHRONIC HA DUE TO NAGS: INITIAL: 6 MOS, RENEWAL: 12 MOS.
<b>Other Criteria</b>	RENEWAL: CHRONIC HA DUE TO NAGS: PATIENT HAS SHOWN CLINICAL IMPROVEMENT.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# CERITINIB

---

## Products Affected

- ZYKADIA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# CERTOLIZUMAB PEGOL

## Products Affected

- CIMZIA (2 SYRINGE)
- CIMZIA SUBCUTANEOUS KIT 2 X 200 MG

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP, OR GENITAL AREA. NON-RADIOGRAPHIC AXIAL SPONDYLOARTHRITIS (NR-AXSPA): 1) C-REACTIVE PROTEIN (CRP) LEVELS ABOVE THE UPPER LIMIT OF NORMAL, OR 2) SACROILIITIS ON MAGNETIC RESONANCE IMAGING (MRI).
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: RHEUMATOID ARTHRITIS (RA), ANKYLOSING SPONDYLITIS (AS), NR-AXSPA, POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. CROHNS DISEASE (CD): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ, RINVOQ, ORENCIA. PSA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: COSENTYX, ENBREL, HUMIRA/CYLTEZO/YUFLYMA, STELARA/SELARSDI/YESINTEK, XELJANZ, RINVOQ, SKYRIZI, TREMFYA, ORENCIA, OTEZLA. PSO: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: COSENTYX, ENBREL,

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>HUMIRA/CYLTEZO/YUFLYMA, STELARA/SELARSDI/YESINTEK, SKYRIZI, TREMFYA, OTEZLA. AS: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: COSENTYX, ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ, RINVOQ. CD: TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING PREFERRED AGENTS: STELARA/SELARSDI/YESINTEK, HUMIRA/CYLTEZO/YUFLYMA, RINVOQ, SKYRIZI, TREMFYA. NR-AXSPA: TRIAL OF OR CONTRAINDICATION TO AN NSAID. PJIA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ IR, ORENCIA, RINVOQ. INITIAL FOR RA, PSA, PSO, AS, CD, PJIA: TRIAL OF OR CONTRAINDICATION TO THE STEP AGENTS IS NOT REQUIRED IF THE PATIENT IS PREGNANT, BREASTFEEDING, OR TRYING TO BECOME PREGNANT. INITIAL/RENEWAL FOR PSA, PSO, AS, CD, NR-AXSPA, PJIA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR SAME INDICATION. RENEWAL FOR RA, PSA, AS, PSO, NR-AXSPA, PJIA: CONTINUES TO BENEFIT FROM MEDICATION.</p>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# CETUXIMAB

---

## Products Affected

- ERBITUX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# CLADRIBINE

## Products Affected

- MAVENCLAD (10 TABS)
- MAVENCLAD (4 TABS)
- MAVENCLAD (5 TABS)
- MAVENCLAD (6 TABS)
- MAVENCLAD (7 TABS)
- MAVENCLAD (8 TABS)
- MAVENCLAD (9 TABS)

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	48 WEEKS.
<b>Other Criteria</b>	RELAPSING FORM OF MULTIPLE SCLEROSIS (MS): HAS NOT RECEIVED A TOTAL OF TWO YEARS OF MAVENCLAD TREATMENT (I.E., TWO YEARLY TREATMENT COURSES OF TWO CYCLES IN EACH).
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# CLOBAZAM-SYMPAZAN

## Products Affected

- SYMPAZAN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	LENNOX-GASTAUT SYNDROME (LGS): THERAPY IS PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	LGS: 1) UNABLE TO TAKE TABLETS OR SUSPENSIONS, AND 2) TRIAL OF OR CONTRAINDICATION TO A FORMULARY VERSION OF CLOBAZAM.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# COBIMETINIB

## Products Affected

- COTELLIC

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# CORTICOTROPIN

## Products Affected

- ACTHAR
- ACTHAR GEL SUBCUTANEOUS PEN-INJECTOR 40 UNIT/0.5ML, 80 UNIT/ML
- CORTROPHIN

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	INITIAL: NOT APPROVED FOR DIAGNOSTIC PURPOSES.
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: ALL FDA APPROVED INDICATIONS EXCEPT INFANTILE SPASMS AND MULTIPLE SCLEROSIS (MS): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST, DERMATOLOGIST, ALLERGIST/IMMUNOLOGIST, OPHTHALMOLOGIST, PULMONOLOGIST OR NEPHROLOGIST.
<b>Coverage Duration</b>	INFANTILE SPASMS AND MS: 28 DAYS. ALL OTHER FDA APPROVED INDICATIONS: INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: ALL FDA APPROVED INDICATIONS EXCEPT INFANTILE SPASMS: TRIAL OF OR CONTRAINDICATION TO INTRAVENOUS (IV) CORTICOSTEROIDS. RENEWAL: ALL FDA APPROVED INDICATIONS EXCEPT INFANTILE SPASMS AND MS: DEMONSTRATED CLINICAL BENEFIT WHILE ON THERAPY AS INDICATED BY SYMPTOM RESOLUTION AND/OR NORMALIZATION OF LABORATORY TESTS. PART B BEFORE PART D STEP THERAPY, APPLIES ONLY TO BENEFICIARIES IN AN MA-PD PLAN.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	Yes

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# CRIZOTINIB CAPSULE

## Products Affected

- XALKORI ORAL CAPSULE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# CRIZOTINIB PELLETS

## Products Affected

- XALKORI ORAL CAPSULE SPRINKLE  
150 MG, 20 MG, 50 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	NON-SMALL CELL LUNG CANCER (NSCLC), ANAPLASTIC LARGE CELL LYMPHOMA (ALCL), INFLAMMATORY MYOFIBROBLASTIC TUMOR (IMT); UNABLE TO SWALLOW CAPSULES.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# DABRAFENIB CAPSULES

---

## Products Affected

- TAFINLAR ORAL CAPSULE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# DABRAFENIB SUSPENSION

---

## Products Affected

- TAFINLAR ORAL TABLET SOLUBLE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	UNABLE TO SWALLOW TAFINLAR CAPSULES.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# DACOMITINIB

---

## Products Affected

- VIZIMPRO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC): NOT ON CONCURRENT THERAPY WITH AN EGFR TYROSINE KINASE-INHIBITOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# DALFAMPRIDINE

## Products Affected

- *dalfampridine er*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	MULTIPLE SCLEROSIS (MS): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL: 3 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	MS: INITIAL: HAS SYMPTOMS OF A WALKING DISABILITY SUCH AS MILD TO MODERATE BILATERAL LOWER EXTREMITY WEAKNESS OR UNILATERAL WEAKNESS PLUS LOWER EXTREMITY OR TRUNCAL ATAXIA. RENEWAL: IMPROVEMENT IN WALKING ABILITY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# DAROLUTAMIDE

## Products Affected

- NUBEQA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS
Other Criteria	INITIAL: NON-METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (NMCRPC): HIGH RISK PROSTATE CANCER (I.E., RAPIDLY INCREASING PROSTATE SPECIFIC ANTIGEN [PSA] LEVELS). NMCRPC, METASTATIC HORMONE-SENSITIVE PROSTATE CANCER (MHSPC): 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG. RENEWAL: NMCRPC, MHSPC: 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GNRH ANALOG.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# DASATINIB

## Products Affected

- *dasatinib oral tablet 100 mg, 140 mg, 20 mg, 50 mg, 70 mg, 80 mg*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	PREVIOUSLY TREATED PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (Ph+ CML): MUTATIONAL ANALYSIS PRIOR TO INITIATION AND DASATINIB IS APPROPRIATE PER NCCN GUIDELINE TABLE FOR TREATMENT RECOMMENDATIONS BASED ON BCR-ABL1 MUTATION PROFILE.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# DATOPOTAMAB DERUXTECAN-DLNK

---

## Products Affected

- DATROWAY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# DECITABINE/CEDAZURIDINE

## Products Affected

- INQOVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# DEFERASIROX

## Products Affected

- *deferasirox granules*
- *deferasirox oral tablet*
- *deferasirox oral tablet soluble 250 mg, 500 mg*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: CHRONIC IRON OVERLOAD DUE TO BLOOD TRANSFUSIONS: SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 1000 MCG/L (AT LEAST TWO LAB VALUES IN THE PREVIOUS THREE MONTHS). CHRONIC IRON OVERLOAD IN NON-TRANSFUSION DEPENDENT THALASSEMIA (NTDT): 1) SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 300 MCG/L (AT LEAST TWO LAB VALUES IN THE PREVIOUS THREE MONTHS), AND 2) LIVER IRON CONCENTRATION (LIC) OF 5 MG FE/G OF DRY LIVER WEIGHT OR GREATER. RENEWAL: CHRONIC IRON OVERLOAD DUE TO BLOOD TRANSFUSIONS: SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 500 MCG/L (AT LEAST TWO LAB VALUES IN THE PREVIOUS THREE MONTHS). NTDT: 1) SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 300 MCG/L (AT LEAST TWO LAB VALUES IN THE PREVIOUS THREE MONTHS) OR 2) LIC OF 3 MG FE/G OF DRY LIVER WEIGHT OR GREATER.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL (CHRONIC IRON OVERLOAD): PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR HEMATOLOGIST/ONCOLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL (CHRONIC IRON OVERLOAD): DEFERASIROX SPRINKLE PACKETS: TRIAL OF OR CONTRAINDICATION TO GENERIC DEFERASIROX ORAL TABLET OR TABLET FOR ORAL SUSPENSION.
<b>Indications</b>	All FDA-approved Indications.

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# DENOSUMAB-XGEVA

## Products Affected

- XGEVA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# DEUTETRABENAZINE

## Products Affected

- AUSTEDO ORAL TABLET 12 MG, 6 MG, 9 MG
- AUSTEDO XR ORAL TABLET EXTENDED RELEASE 24 HOUR 12 MG, 18 MG, 24 MG, 30 MG, 36 MG, 42 MG, 48 MG, 6 MG
- AUSTEDO XR PATIENT TITRATION

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	HUNTINGTON DISEASE: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST OR MOVEMENT DISORDER SPECIALIST. TARDIVE DYSKINESIA: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST, PSYCHIATRIST, OR MOVEMENT DISORDER SPECIALIST.
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	TARDIVE DYSKINESIA: HISTORY OF USING AGENTS THAT CAUSE TARDIVE DYSKINESIA.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# DEXTROMETHORPHAN QUINIDINE

## Products Affected

- NUEDEXTA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All Medically-accepted Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# DICLOFENAC - FLECTOR

## Products Affected

- *diclofenac epolamine external*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All Medically-accepted Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# DICLOFENAC TOPICAL SOLUTION

## Products Affected

- *diclofenac sodium external solution 2 %*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 MONTHS
Other Criteria	OSTEOARTHRITIS OF THE KNEE: TRIAL OF OR CONTRAINDICATION TO A FORMULARY VERSION OF DICLOFENAC SODIUM 1% TOPICAL GEL AND A FORMULARY VERSION OF DICLOFENAC SODIUM 1.5% TOPICAL DROPS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# DIMETHYL FUMARATE

## Products Affected

- *dimethyl fumarate oral capsule delayed release 120 mg, 240 mg*
- *dimethyl fumarate starter pack oral capsule delayed release therapy pack*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# DIROXIMEL FUMARATE

---

## Products Affected

- VUMERITY

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# DOSTARLIMAB-GXLY

---

## Products Affected

- JEMPERLI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# DRONABINOL CAPSULE

## Products Affected

- *dronabinol*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 MONTHS
Other Criteria	NAUSEA AND VOMITING ASSOCIATED WITH CANCER CHEMOTHERAPY: TRIAL OF OR CONTRAINDICATION TO ONE ANTIEMETIC THERAPY. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D FOR THE INDICATION OF NAUSEA AND VOMITING ASSOCIATED WITH CANCER CHEMOTHERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# DROXIDOPA

## Products Affected

- *droxidopa*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	NEUROGENIC ORTHOSTATIC HYPOTENSION (NOH): INITIAL: 1) BASELINE BLOOD PRESSURE READINGS WHILE THE PATIENT IS SITTING AND ALSO WITHIN 3 MINUTES OF STANDING FROM A SUPINE POSITION. 2) A DECREASE OF AT LEAST 20 MMHG IN SYSTOLIC BLOOD PRESSURE OR 10 MMHG DIASTOLIC BLOOD PRESSURE WITHIN THREE MINUTES AFTER STANDING FROM A SITTING POSITION.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	NOH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST OR CARDIOLOGIST.
<b>Coverage Duration</b>	INITIAL: 3 MONTHS RENEWAL: 12 MONTHS
<b>Other Criteria</b>	NOH: RENEWAL: CONTINUES TO BENEFIT FROM THE MEDICATION.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# DUPILUMAB

## Products Affected

- DUPIXENT SUBCUTANEOUS SOLUTION AUTO-INJECTOR
- DUPIXENT SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: EOSINOPHILIC ASTHMA: BLOOD EOSINOPHIL LEVEL OF 150 TO 1500 CELLS/MCL WITHIN THE PAST 12 MONTHS. EOSINOPHILIC ESOPHAGITIS (EOE): DIAGNOSIS CONFIRMED BY ESOPHAGOGASTRODUODENOSCOPY (EGD) WITH BIOPSY. ATOPIC DERMATITIS (AD): AD COVERING AT LEAST 10 PERCENT OF BODY SURFACE AREA OR AD AFFECTING THE FACE, HEAD, NECK, HANDS, FEET, GROIN, OR INTERTRIGINOUS AREAS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: AD, PRURIGO NODULARIS (PN): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST, ALLERGIST OR IMMUNOLOGIST. ASTHMA: PRESCRIBED BY OR IN CONSULTATION WITH A PHYSICIAN SPECIALIZING IN ALLERGY OR PULMONARY MEDICINE. CRSWNP: PRESCRIBED BY OR IN CONSULTATION WITH AN OTOLARYNGOLOGIST, ALLERGIST OR IMMUNOLOGIST. EOE: PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST, ALLERGIST, OR IMMUNOLOGIST. EOSINOPHILIC COPD: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST.
<b>Coverage Duration</b>	INITIAL: AD, CRSWNP, EOE, PN: 6 MOS, ASTHMA, COPD: 12 MOS. RENEWAL: ALL INDICATIONS: 12 MOS.
<b>Other Criteria</b>	INITIAL: AD: 1) INTRACTABLE PRURITUS OR CRACKING/OOZING/BLEEDING OF AFFECTED SKIN, 2) TRIAL OF OR CONTRAINDICATION TO ONE TOPICAL (CORTICOSTEROID, CALCINEURIN INHIBITOR, PDE4 INHIBITOR, OR JAK INHIBITOR), AND 3) NO CONCURRENT USE WITH OTHER SYSTEMIC BIOLOGICS OR JAK INHIBITORS FOR AD. ASTHMA: 1) CONCURRENT THERAPY WITH A MEDIUM,

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>HIGH-DOSE OR MAXIMALLY-TOLERATED DOSE OF AN INHALED CORTICOSTEROID (ICS) AND ONE OTHER MAINTENANCE MEDICATION, 2) ONE ASTHMA EXACERBATION REQUIRING SYSTEMIC CORTICOSTEROID BURST LASTING 3 OR MORE DAYS WITHIN THE PAST 12 MONTHS, OR ONE SERIOUS EXACERBATION REQUIRING HOSPITALIZATION OR ER VISIT WITHIN THE PAST 12 MONTHS, OR POOR SYMPTOM CONTROL DESPITE CURRENT THERAPY AS EVIDENCED BY AT LEAST THREE OF THE FOLLOWING WITHIN THE PAST 4 WEEKS: (A) DAYTIME ASTHMA SYMPTOMS MORE THAN TWICE/WEEK, (B) ANY NIGHT WAKING DUE TO ASTHMA, (C) SABA RELIEVER FOR SYMPTOMS MORE THAN TWICE/WEEK, (D) ANY ACTIVITY LIMITATION DUE TO ASTHMA, AND 3) NO CONCURRENT USE WITH XOLAIR, TEZSPIRE, OR OTHER ANTI-IL5 BIOLOGICS WHEN USED FOR ASTHMA. CHRONIC RHINOSINUSITIS WITH NASAL POLYPS (CRSWNP): 1) A 56 DAY TRIAL OF ONE TOPICAL NASAL CORTICOSTEROID, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR AN AUTOIMMUNE INDICATION. PN: 1) CHRONIC PRURITIS (ITCH MORE THAN 6 WEEKS), MULTIPLE PRURIGINOUS LESIONS, AND HISTORY OR SIGN OF A PROLONGED SCRATCHING BEHAVIOR, AND 2) TRIAL OF OR CONTRAINDICATION TO ONE TOPICAL (CORTICOSTEROID OR CALCIPOTRIOL). EOSINOPHILIC COPD: 1) USED IN COMBINATION WITH A LAMA/LABA/ICS, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR EOSINOPHILIC COPD. RENEWAL: AD: 1) IMPROVEMENT WHILE ON THERAPY, AND 2) NO CONCURRENT USE WITH OTHER SYSTEMIC BIOLOGICS OR JAK INHIBITORS FOR AD. EOE: IMPROVEMENT WHILE ON THERAPY. CRSWNP: 1) IMPROVEMENT WHILE ON THERAPY, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AN AUTOIMMUNE INDICATION. ASTHMA: 1) NO CONCURRENT USE WITH XOLAIR, TEZSPIRE, OR OTHER ANTI-IL5 BIOLOGICS FOR ASTHMA, 2) CONTINUED USE OF ICS AND ONE OTHER MAINTENANCE MEDICATION, AND 3) CLINICAL RESPONSE AS EVIDENCED BY: (A) REDUCTION IN ASTHMA EXACERBATIONS FROM BASELINE, (B) DECREASED UTILIZATION OF RESCUE MEDICATIONS, (C) INCREASE IN PERCENT PREDICTED FEV1</p>

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	FROM PRETREATMENT BASELINE, OR (D) REDUCTION IN SEVERITY OR FREQUENCY OF ASTHMA-RELATED SYMPTOMS. PN: IMPROVEMENT OR REDUCTION OF PRURITIS OR PRURIGINOUS LESIONS. EOSINOPHILIC COPD: 1) USED IN COMBINATION WITH A LAMA/LABA/ICS, 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR EOSINOPHILIC COPD, AND 3) CLINICAL RESPONSE AS EVIDENCED BY (A) REDUCTION IN COPD EXACERBATIONS FROM BASELINE, (B) REDUCTION IN SEVERITY OR FREQUENCY OF COPD-RELATED SYMPTOMS, OR (C) INCREASE IN FEV1 OF AT LEAST 5 PERCENT FROM PRETREATMENT BASELINE.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# DUVELISIB

---

## Products Affected

- COPIKTRA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# EFLORNITHINE

---

## Products Affected

- IWILFIN

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	24 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ELACESTRANT

## Products Affected

- ORSERDU ORAL TABLET 345 MG, 86 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ELAGOLIX

## Products Affected

- ORLISSA ORAL TABLET 150 MG, 200 MG

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS: INITIAL: DIAGNOSIS IS CONFIRMED VIA SURGICAL OR DIRECT VISUALIZATION (E.G., PELVIC ULTRASOUND) OR HISTOPATHOLOGICAL CONFIRMATION (E.G., LAPAROSCOPY OR LAPAROTOMY) IN THE LAST 10 YEARS.
<b>Age Restrictions</b>	MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS: INITIAL: 18 YEARS OF AGE OR OLDER.
<b>Prescriber Restrictions</b>	MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH AN OBSTETRICIAN/GYNECOLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS
<b>Other Criteria</b>	MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS: INITIAL: 1) NO CONCURRENT USE WITH ANOTHER GNRH-MODULATING AGENT, AND 2) TRIAL OF OR CONTRAINDICATION TO AN NSAID AND A PROGESTIN-CONTAINING PREPARATION. RENEWAL: 1) IMPROVEMENT IN PAIN ASSOCIATED WITH ENDOMETRIOSIS WHILE ON THERAPY, AND 2) NO CONCURRENT USE WITH ANOTHER GNRH-MODULATING AGENT
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# ELRANATAMAB-BCMM

## Products Affected

- ELREXFIO SUBCUTANEOUS SOLUTION 44 MG/1.1ML, 76 MG/1.9ML

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	RELAPSED OR REFRACTORY MULTIPLE MYELOMA: RENEWAL: 1) HAS RECEIVED AT LEAST 24 WEEKS OF TREATMENT WITH ELREXFIO, AND 2) HAS RESPONDED TO TREATMENT (PARTIAL RESPONSE OR BETTER), AND HAS MAINTAINED THIS RESPONSE FOR AT LEAST 2 MONTHS. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# ELTROMBOPAG - ALVAIZ

## Products Affected

- ALVAIZ

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	PERSISTENT OR CHRONIC IMMUNE THROMBOCYTOPENIA (ITP): INITIAL: 1) PLATELET COUNT IS LESS THAN 30 X 10 <sup>9</sup> /L FROM AT LEAST 2 SEPARATE LABS IN THE LAST 3 MONTHS, OR 2) PLATELET COUNT IS LESS THAN 50 X 10 <sup>9</sup> /L FROM AT LEAST 2 SEPARATE LABS IN THE LAST 3 MONTHS AND HAD A PRIOR BLEEDING EVENT.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: ITP: PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR IMMUNOLOGIST.
<b>Coverage Duration</b>	ITP: INITIAL: 6 MO, RENEWAL: 12 MO. HEPATITIS C, SEVERE APLASTIC ANEMIA: 12 MO.
<b>Other Criteria</b>	INITIAL: ITP: 1) TRIAL OF OR CONTRAINDICATION TO ONE CORTICOSTEROID OR IMMUNOGLOBULIN, OR AN INSUFFICIENT RESPONSE TO SPLENECTOMY, AND 2) NO CONCURRENT USE WITH OTHER THROMBOPOIETIN RECEPTOR AGONISTS (TPO-RAS) OR SPLEEN TYROSINE KINASE (SYK) INHIBITOR. RENEWAL: ITP: 1) IMPROVEMENT IN PLATELET COUNT FROM BASELINE OR REDUCTION IN BLEEDING EVENTS, AND 2) NO CONCURRENT USE WITH OTHER TPO-RAS OR SYK INHIBITOR.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# ELTROMBOPAG - PROMACTA

## Products Affected

- PROMACTA ORAL PACKET 12.5 MG, 25 MG
- PROMACTA ORAL TABLET 12.5 MG, 25 MG, 50 MG, 75 MG

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	PERSISTENT OR CHRONIC IMMUNE THROMBOCYTOPENIA (ITP): INITIAL: 1) PLATELET COUNT OF LESS THAN $30 \times 10^9/L$ FROM AT LEAST 2 SEPARATE LAB TESTS IN THE LAST 3 MONTHS, OR 2) PLATELET COUNT OF LESS THAN $50 \times 10^9/L$ FROM AT LEAST 2 SEPARATE LAB TESTS IN THE LAST 3 MONTHS AND A PRIOR BLEEDING EVENT.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: ITP: PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR IMMUNOLOGIST.
<b>Coverage Duration</b>	ITP: INITIAL: 6 MO, RENEWAL: 12 MO. HEPATITIS C, SEVERE APLASTIC ANEMIA: 12 MO.
<b>Other Criteria</b>	INITIAL: ITP: 1) TRIAL OF OR CONTRAINDICATION TO ONE CORTICOSTEROID OR IMMUNOGLOBULIN, OR HAD AN INSUFFICIENT RESPONSE TO SPLENECTOMY, AND 2) NO CONCURRENT USE WITH OTHER THROMBOPOIETIN RECEPTOR AGONISTS (TPO-RAS) OR SPLEEN TYROSINE KINASE (SYK) INHIBITOR. ALL INDICATIONS: APPROVAL FOR PROMACTA ORAL SUSPENSION PACKETS REQUIRES A TRIAL OF PROMACTA TABLET OR PATIENT IS UNABLE TOLERATE TABLET FORMULATION. RENEWAL: ITP: 1) IMPROVEMENT IN PLATELET COUNTS FROM BASELINE OR REDUCTION IN BLEEDING EVENTS, AND 2) NO CONCURRENT USE WITH OTHER TPO-RAS OR SYK INHIBITOR.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ENASIDENIB

---

## Products Affected

- IDHIFA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ENCORAFENIB

---

## Products Affected

- BRAFTOVI ORAL CAPSULE 75 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ENTRECTINIB CAPSULES

## Products Affected

- ROZLYTREK ORAL CAPSULE 100 MG, 200 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# ENTRECTINIB PELLETS

## Products Affected

- ROZLYTREK ORAL PACKET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), SOLID TUMORS: 1) TRIAL OF OR CONTRAINDICATION TO ROZLYTREK CAPSULES MADE INTO AN ORAL SUSPENSION, AND 2) DIFFICULTY OR UNABLE TO SWALLOW CAPSULES.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# ENZALUTAMIDE

## Products Affected

- XTANDI ORAL CAPSULE
- XTANDI ORAL TABLET 40 MG, 80 MG

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	INITIAL: ALL INDICATIONS: 12 MONTHS. RENEWAL: MCRPC, NMCRPC, MCSPC: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: NON-METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (NMCRPC): HIGH RISK PROSTATE CANCER (I.E. RAPIDLY INCREASING PROSTATE SPECIFIC ANTIGEN [PSA] LEVELS). NON-METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (NMCSPC): HIGH RISK FOR METASTASIS (I.E. PSA DOUBLING TIME OF 9 MONTHS OR LESS). METASTATIC CRPC (MCRPC), NMCRPC, METASTATIC CSPC (MCSPC), NMCSPC : 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG. RENEWAL: MCRPC, NMCRPC, MCSPC: 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GNRH ANALOG.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# EPCORITAMAB-BYSP

## Products Affected

- EPKINLY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# EPOETIN ALFA - PROCRIT

## Products Affected

- PROCRIT INJECTION SOLUTION                      UNIT/ML, 3000 UNIT/ML, 4000  
10000 UNIT/ML, 2000 UNIT/ML, 20000                      UNIT/ML, 40000 UNIT/ML

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: CHRONIC KIDNEY DISEASE (CKD), ANEMIA RELATED TO ZIDOVUDINE, OR CANCER CHEMOTHERAPY: HEMOGLOBIN LEVEL IS LESS THAN 10G/DL. ELECTIVE, NON-CARDIAC, NON-VASCULAR SURGERY: HEMOGLOBIN LEVEL IS LESS THAN 13G/DL. RENEWAL: 1) CKD IN ADULTS NOT ON DIALYSIS: (A) HEMOGLOBIN LEVEL IS LESS THAN 10G/DL, OR (B) HEMOGLOBIN LEVEL HAS REACHED 10G/DL AND THE DOSE IS BEING OR HAS BEEN REDUCED/INTERRUPTED TO DECREASE THE NEED FOR BLOOD TRANSFUSIONS. 2) CKD IN PEDIATRIC PATIENTS: (A) HEMOGLOBIN LEVEL IS LESS THAN 10G/DL, OR (B) HEMOGLOBIN LEVEL HAS APPROACHED OR EXCEEDS 12G/DL AND THE DOSE IS BEING OR HAS BEEN REDUCED/INTERRUPTED TO DECREASE THE NEED FOR BLOOD TRANSFUSIONS. 3) ANEMIA RELATED TO ZIDOVUDINE: HEMOGLOBIN LEVEL BETWEEN 10G/DL AND 12G/DL. 4) CANCER CHEMOTHERAPY: (A) HEMOGLOBIN LEVEL IS LESS THAN 10 G/DL, OR (B) HEMOGLOBIN LEVEL DOES NOT EXCEED A LEVEL NEEDED TO AVOID RBC TRANSFUSION.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	ANEMIA FROM CHEMO/CKD WITHOUT DIALYSIS/ZIDOVUDINE: INITIAL/RENEWAL: 12 MONTHS. SURGERY: 1 MONTH.
<b>Other Criteria</b>	RENEWAL: CKD: NOT RECEIVING DIALYSIS TREATMENT. THIS DRUG MAY BE EITHER BUNDLED WITH AND COVERED UNDER END STAGE RENAL DISEASE DIALYSIS RELATED SERVICES OR COVERED UNDER MEDICARE D DEPENDING UPON THE CIRCUMSTANCES.

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# EPOETIN ALFA-EPBX

## Products Affected

- RETACRIT INJECTION SOLUTION                      UNIT/ML, 4000 UNIT/ML, 40000  
10000 UNIT/ML, 10000 UNIT/ML(1ML),            UNIT/ML  
2000 UNIT/ML, 20000 UNIT/ML, 3000

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: CHRONIC KIDNEY DISEASE (CKD), ANEMIA RELATED TO ZIDOVUDINE, OR CANCER CHEMOTHERAPY: HEMOGLOBIN LEVEL IS LESS THAN 10G/DL. ELECTIVE, NON-CARDIAC, NON-VASCULAR SURGERY: HEMOGLOBIN LEVEL IS LESS THAN 13G/DL. RENEWAL: 1) CKD IN ADULTS NOT ON DIALYSIS: (A) HEMOGLOBIN LEVEL IS LESS THAN 10G/DL, OR (B) HEMOGLOBIN LEVEL HAS REACHED 10G/DL AND THE DOSE IS BEING OR HAS BEEN REDUCED/INTERRUPTED TO DECREASE THE NEED FOR BLOOD TRANSFUSIONS. 2) CKD IN PEDIATRIC PATIENTS: (A) HEMOGLOBIN LEVEL IS LESS THAN 10G/DL, OR (B) HEMOGLOBIN LEVEL HAS APPROACHED OR EXCEEDS 12G/DL AND THE DOSE IS BEING OR HAS BEEN REDUCED/INTERRUPTED TO DECREASE THE NEED FOR BLOOD TRANSFUSIONS. 3) ANEMIA RELATED TO ZIDOVUDINE: HEMOGLOBIN LEVEL BETWEEN 10G/DL AND 12G/DL. 4) CANCER CHEMOTHERAPY: (A) HEMOGLOBIN LEVEL IS LESS THAN 10 G/DL, OR (B) HEMOGLOBIN LEVEL DOES NOT EXCEED A LEVEL NEEDED TO AVOID RBC TRANSFUSION.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	ANEMIA FROM CHEMO/CKD WITHOUT DIALYSIS/ZIDOVUDINE: INITIAL/RENEWAL: 12 MONTHS. SURGERY: 1 MONTH.
<b>Other Criteria</b>	RENEWAL: CKD: NOT RECEIVING DIALYSIS TREATMENT. THIS DRUG MAY BE EITHER BUNDLED WITH AND COVERED UNDER END STAGE RENAL DISEASE DIALYSIS RELATED SERVICES OR COVERED UNDER MEDICARE D DEPENDING UPON THE CIRCUMSTANCES.

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ERDAFITINIB

## Products Affected

- BALVERSA ORAL TABLET 3 MG, 4 MG, 5 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ERENUMAB - AOOE

## Products Affected

- AIMOVIG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	MIGRAINE PREVENTION: INITIAL: 1) TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING PREVENTIVE MIGRAINE TREATMENTS: DIVALPROEX SODIUM, TOPIRAMATE, PROPRANOLOL, TIMOLOL, AND 2) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR MIGRAINE PREVENTION. RENEWAL: 1) REDUCTION IN MIGRAINE OR HEADACHE FREQUENCY, MIGRAINE SEVERITY, OR MIGRAINE DURATION WITH THERAPY, AND 2) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR MIGRAINE PREVENTION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# ERLOTINIB

## Products Affected

- *erlotinib hcl oral tablet 100 mg, 150 mg, 25 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EGFR MUTATION: NOT ON CONCURRENT THERAPY WITH AN EGFR TYROSINE KINASE INHIBITOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ESKETAMINE

## Products Affected

- SPRAVATO (56 MG DOSE)
- SPRAVATO (84 MG DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: TREATMENT-RESISTANT DEPRESSION (TRD), MAJOR DEPRESSIVE DISORDER (MDD): PRESCRIBED BY OR IN CONSULTATION WITH A PSYCHIATRIST.
Coverage Duration	INITIAL: TRD: 3 MONTHS. MDD: 4 WEEKS. RENEWAL: TRD, MDD: 12 MONTHS.
Other Criteria	INITIAL: TRD, MDD: 1) NON-PSYCHOTIC, UNIPOLAR DEPRESSION, AND 2) NO ACTIVE SUBSTANCE ABUSE. RENEWAL: TRD, MDD: DEMONSTRATED CLINICAL BENEFIT (IMPROVEMENT IN DEPRESSION) COMPARED TO BASELINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# ETANERCEPT

## Products Affected

- ENBREL MINI
- ENBREL SUBCUTANEOUS SOLUTION 25 MG/0.5ML
- ENBREL SUBCUTANEOUS SOLUTION PREFILLED SYRINGE
- ENBREL SUBCUTANEOUS SOLUTION RECONSTITUTED
- ENBREL SURECLICK SUBCUTANEOUS SOLUTION AUTO-INJECTOR

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP, OR GENITAL AREA.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: RHEUMATOID ARTHRITIS (RA), POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA), ANKYLOSING SPONDYLITIS (AS): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. PJIA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PJIA. PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. AS: 1) TRIAL OF OR CONTRAINDICATION TO AN NSAID, AND 2) NO CONCURRENT

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. PSO: 1) ONE OF THE FOLLOWING: (A) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY) FOR THE TREATMENT OF PSO, (B) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA FOR THE TREATMENT OF PSO, OR (C) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. RENEWAL: RA: CONTINUES TO BENEFIT FROM THE MEDICATION. PJIA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PJIA. PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. AS: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. PSO: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO.</p>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# EVEROLIMUS-AFINITOR

## Products Affected

- *everolimus oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg*
- *torpenz oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# EVEROLIMUS-AFINITOR DISPERZ

---

## Products Affected

- *everolimus oral tablet soluble*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# FECAL MICROBIOTA CAPSULE

## Products Affected

- VOWST

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	30 DAYS
Other Criteria	CLOSTRIDIODES DIFFICILE INFECTION (CDI): 1) HAS NOT PREVIOUSLY RECEIVED VOWST: COMPLETION OF ANTIBIOTIC TREATMENT FOR RECURRENT CDI (AT LEAST 3 CDI EPISODES), OR 2) PREVIOUSLY RECEIVED VOWST: (A) TREATMENT FAILURE (DEFINED AS THE PRESENCE OF CDI DIARRHEA WITHIN 8 WEEKS OF FIRST DOSE OF VOWST AND A POSITIVE STOOL TEST FOR C. DIFFICILE), AND (B) HAS NOT RECEIVED MORE THAN ONE TREATMENT COURSE OF VOWST WHICH WAS AT LEAST 12 DAYS AND NOT MORE THAN 8 WEEKS PRIOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# FEDRATINIB

## Products Affected

- INREBIC

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	MYELOFIBROSIS: INITIAL: TRIAL OF OR CONTRAINDICATION TO JAKAFI (RUXOLITINIB). RENEWAL: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# FENFLURAMINE

## Products Affected

- FINTEPLA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: DRAVET SYNDROME, LENNOX-GASTAUT SYNDROME (LGS): PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	DRAVET SYNDROME: INITIAL/RENEWAL: 12 MONTHS. LGS: 12 MONTHS.
Other Criteria	INITIAL: LGS: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING ANTIEPILEPTIC MEDICATIONS: RUFINAMIDE, FELBAMATE, CLOBAZAM, TOPIRAMATE, LAMOTRIGINE, CLONAZEPAM. RENEWAL: DRAVET SYNDROME: PATIENT HAS SHOWN CONTINUED CLINICAL BENEFIT (E.G. REDUCTION OF SEIZURES, REDUCED LENGTH OF SEIZURES, SEIZURE CONTROL MAINTAINED).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# FENTANYL CITRATE

## Products Affected

- *fentanyl citrate buccal lozenge on a handle*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	CANCER RELATED PAIN: 1) CURRENTLY ON A MAINTENANCE DOSE OF CONTROLLED-RELEASE OPIOID PAIN MEDICATION, AND 2) TRIAL OF OR CONTRAINDICATION TO AT LEAST ONE IMMEDIATE-RELEASE ORAL OPIOID PAIN AGENT OR PATIENT HAS DIFFICULTY SWALLOWING TABLETS/CAPSULES.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# FEZOLINETANT

## Products Affected

- VEOZAH

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	MENOPAUSAL VASOMOTOR SYMPTOMS (VMS): INITIAL: 1) EXPERIENCES 7 OR MORE HOT FLASHES PER DAY, AND 2) TRIAL OF OR CONTRAINDICATION TO HORMONAL THERAPY (E.G., ESTRADIOL TRANSDERMAL PATCH, ORAL CONJUGATED ESTROGENS). RENEWAL: 1) CONTINUED NEED FOR VMS TREATMENT (I.E., PERSISTENT HOT FLASHES), AND 2) REDUCTION IN VMS FREQUENCY OR SEVERITY DUE TO VEOZAH TREATMENT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# FILGRASTIM-AAFI

## Products Affected

- NIVESTYM

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# FINERENONE

---

## Products Affected

- KERENDIA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# FINGOLIMOD

---

## Products Affected

- *fingolimod hcl*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# FOSCARBIDOPA-FOSLEVODOPA

## Products Affected

- VYALEV SUBCUTANEOUS SOLUTION 12-240 MG/ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PARKINSONS DISEASE (PD): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL: 3 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	PD: INITIAL: 1) RESPONSIVE TO LEVODOPA, 2) CURRENT REGIMEN INCLUDES AT LEAST 400 MG/DAY OF LEVODOPA, AND 3) MOTOR SYMPTOMS ARE CURRENTLY UNCONTROLLED (DEFINED AS AN AVERAGE OFF TIME OF AT LEAST 2.5 HOURS/DAY OVER 3 CONSECUTIVE DAYS WITH A MINIMUM OF 2 HOURS EACH DAY). RENEWAL: IMPROVEMENT IN MOTOR SYMPTOMS WHILE ON THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# FREMANEZUMAB-VFRM

## Products Affected

- AJOVY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS.
Other Criteria	MIGRAINE PREVENTION: INITIAL: 1) TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING PREVENTIVE MIGRAINE TREATMENTS: DIVALPROEX SODIUM, TOPIRAMATE, PROPRANOLOL, TIMOLOL, AND 2) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR MIGRAINE PREVENTION. RENEWAL: 1) REDUCTION IN MIGRAINE OR HEADACHE FREQUENCY, MIGRAINE SEVERITY, OR MIGRAINE DURATION WITH THERAPY, AND 2) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR MIGRAINE PREVENTION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# FRUQUINTINIB

## Products Affected

- FRUZAQLA ORAL CAPSULE 1 MG, 5 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# FUTIBATINIB

## Products Affected

- LYTGOBI (12 MG DAILY DOSE)
- LYTGOBI (16 MG DAILY DOSE)
- LYTGOBI (20 MG DAILY DOSE)

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INTRAHEPATIC CHOLANGIOCARCINOMA (ICCA): COMPLETE A COMPREHENSIVE OPHTHALMOLOGICAL EXAMINATION, INCLUDING OPTICAL COHERENCE TOMOGRAPHY (OCT), PRIOR TO THE INITIATION OF THERAPY AND AT THE RECOMMENDED SCHEDULED INTERVALS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# GALCANEZUMAB-GNLM

## Products Affected

- EMGALITY
- EMGALITY (300 MG DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: MIGRAINE PREVENTION: 6 MOS. EPISODIC CLUSTER HEADACHE: 3 MOS. RENEWAL (ALL): 12 MOS.
Other Criteria	INITIAL: MIGRAINE PREVENTION: 1) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR MIGRAINE PREVENTION, AND 2) TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING PREVENTIVE MIGRAINE TREATMENTS: DIVALPROEX SODIUM, TOPIRAMATE, PROPRANOLOL, TIMOLOL. RENEWAL: MIGRAINE PREVENTION: 1) REDUCTION IN MIGRAINE OR HEADACHE FREQUENCY, MIGRAINE SEVERITY, OR MIGRAINE DURATION WITH THERAPY, AND 2) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR MIGRAINE PREVENTION. EPISODIC CLUSTER HEADACHE: IMPROVEMENT IN EPISODIC CLUSTER HEADACHE FREQUENCY AS COMPARED TO BASELINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# GANAXOLONE

---

## Products Affected

- ZTALMY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# GEFITINIB

## Products Affected

- *gefitinib*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EGFR MUTATION: NOT ON CONCURRENT THERAPY WITH AN EGFR TYROSINE KINASE INHIBITOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# GILTERITINIB

---

## Products Affected

- XOSPATA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# GLASDEGIB

---

## Products Affected

- DAURISMO ORAL TABLET 100 MG,  
25 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# GLATIRAMER

## Products Affected

- *glatiramer acetate subcutaneous solution prefilled syringe 20 mg/ml, 40 mg/ml*
- *glatopa subcutaneous solution prefilled syringe 20 mg/ml, 40 mg/ml*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# GLP1-DULAGLUTIDE

## Products Affected

- TRULICITY SUBCUTANEOUS SOLUTION AUTO-INJECTOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All Medically-accepted Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# GLP1-SEMAGLUTIDE

## Products Affected

- OZEMPIC (0.25 OR 0.5 MG/DOSE)
- OZEMPIC (1 MG/DOSE)
- OZEMPIC (2 MG/DOSE)
- RYBELSUS
- RYBELSUS (FORMULATION R2)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All Medically-accepted Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# GLP1-TIRZEPATIDE

## Products Affected

- MOUNJARO SUBCUTANEOUS SOLUTION AUTO-INJECTOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All Medically-accepted Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# GOSERELIN

## Products Affected

- ZOLADEX

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	ENDOMETRIOSIS: DIAGNOSIS IS CONFIRMED VIA SURGICAL OR DIRECT VISUALIZATION (E.G., PELVIC ULTRASOUND) OR HISTOPATHOLOGICAL CONFIRMATION (E.G., LAPAROSCOPY OR LAPAROTOMY) IN THE LAST 10 YEARS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	ENDOMETRIOSIS: PRESCRIBED BY OR IN CONSULTATION WITH AN OBSTETRICIAN/GYNECOLOGIST.
<b>Coverage Duration</b>	STAGE B2-C PROSTATIC CARCINOMA: 4 MOS. ENDOMETRIOSIS: 6 MOS PER LIFETIME. ALL OTHERS: 12 MONTHS.
<b>Other Criteria</b>	ENDOMETRIOSIS: 1) NO CONCURRENT USE WITH ANOTHER GNRH-MODULATING AGENT, 2) TRIAL OF OR CONTRAINDICATION TO NSAID AND PROGESTIN-CONTAINING PREPARATION, AND 3) HAS NOT RECEIVED A TOTAL OF 6 MONTHS OF TREATMENT PER LIFETIME. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# GUSELKUMAB

## Products Affected

- TREMFYA INTRAVENOUS
- TREMFYA ONE-PRESS
- TREMFYA PEN SUBCUTANEOUS SOLUTION AUTO-INJECTOR 200 MG/2ML
- TREMFYA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP, OR GENITAL AREA.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST. ULCERATIVE COLITIS (UC), CROHNS DISEASE (CD): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
<b>Other Criteria</b>	INITIAL: PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PSA. PSO: 1) ONE OF THE FOLLOWING: (A) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY) FOR THE TREATMENT OF PSO, (B) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA FOR THE TREATMENT OF PSO, OR (C) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. UC: NO

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. RENEWAL: PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. PSO: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD.</p>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# IBRUTINIB

---

## Products Affected

- IMBRUVICA ORAL CAPSULE 140 MG, 70 MG
- IMBRUVICA ORAL SUSPENSION
- IMBRUVICA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ICATIBANT

## Products Affected

- *icatibant acetate*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	HEREDITARY ANGIOEDEMA (HAE): DIAGNOSIS CONFIRMED BY COMPLEMENT TESTING.
Age Restrictions	
Prescriber Restrictions	HAE: PRESCRIBED BY OR IN CONSULTATION WITH AN ALLERGIST, IMMUNOLOGIST, OR HEMATOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	HAE: NO CONCURRENT USE WITH OTHER MEDICATIONS FOR TREATMENT OF ACUTE HAE ATTACKS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# IDELALISIB

---

## Products Affected

- ZYDELIG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# IMATINIB

## Products Affected

- *imatinib mesylate oral tablet 100 mg, 400 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ADJUVANT GASTROINTESTINAL STROMAL TUMOR TREATMENT: 36 MONTHS. ALL OTHER DIAGNOSES: 12 MONTHS.
Other Criteria	PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA: PATIENT HAS NOT RECEIVED A PREVIOUS TREATMENT WITH ANOTHER TYROSINE KINASE INHIBITOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# IMATINIB SOLUTION

## Products Affected

- IMKELDI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ADJUVANT GASTROINTESTINAL STROMAL TUMOR TREATMENT: 36 MONTHS. ALL OTHER DIAGNOSES: 12 MONTHS.
Other Criteria	PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA: PATIENT HAS NOT RECEIVED A PREVIOUS TREATMENT WITH ANOTHER TYROSINE KINASE INHIBITOR. ALL INDICATIONS: UNABLE TO SWALLOW GENERIC IMATINIB TABLETS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# IMETELSTAT

---

## Products Affected

- RYTELO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# INAVOLISIB

## Products Affected

- ITOVEBI ORAL TABLET 3 MG, 9 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# INFLIXIMAB

## Products Affected

- *infliximab*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP, OR FACE.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: RHEUMATOID ARTHRITIS (RA), ANKYLOSING SPONDYLITIS (AS): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ, RINVOQ, ORENCIA. PSA: 1) TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: COSENTYX, ENBREL, HUMIRA/CYLTEZO/YUFLYMA, STELARA/SELARSDI/YESINTEK, XELJANZ, RINVOQ, SKYRIZI, TREMFYA, ORENCIA, OTEZLA, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PSA. PSO: 1) TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: COSENTYX, ENBREL, HUMIRA/CYLTEZO/YUFLYMA, STELARA/SELARSDI/YESINTEK, SKYRIZI, TREMFYA, OTEZLA, AND 2) NO CONCURRENT USE

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. AS: 1) TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: COSENTYX, ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ, RINVOQ, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. MODERATE TO SEVERE CD: 1) TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS, WHERE AGES ALIGN: STELARA/SELARSDI/YESINTEK, HUMIRA/CYLTEZO/YUFLYMA, RINVOQ, SKYRIZI, TREMFYA, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: 1) TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS, WHERE AGES ALIGN: STELARA/SELARSDI/YESINTEK, XELJANZ, HUMIRA/CYLTEZO/YUFLYMA, RINVOQ, SKYRIZI, TREMFYA, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC. RENEWAL: RA: CONTINUES TO BENEFIT FROM THE MEDICATION. PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. PSO: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. AS: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. MODERATE TO SEVERE CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.</p>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# INSULIN SUPPLIES PAYMENT DETERMINATION

---

## Products Affected

- ADVOCATE INSULIN SYRINGE 29G X 1/2" 0.3 ML
- ADVOCATE INSULIN SYRINGE 29G X 1/2" 0.5 ML
- ADVOCATE INSULIN SYRINGE 29G X 1/2" 1 ML
- ADVOCATE INSULIN SYRINGE 30G X 5/16" 0.3 ML
- ADVOCATE INSULIN SYRINGE 30G X 5/16" 0.5 ML
- ADVOCATE INSULIN SYRINGE 30G X 5/16" 1 ML
- ADVOCATE INSULIN SYRINGE 31G X 5/16" 0.3 ML
- ADVOCATE INSULIN SYRINGE 31G X 5/16" 0.5 ML
- ADVOCATE INSULIN SYRINGE 31G X 5/16" 1 ML
- ALCOHOL PREP PAD
- ALCOHOL PREP PAD 70 %
- ALCOHOL PREP PADS PAD 70 %
- ALCOHOL SWABS PAD
- ALCOHOL SWABS PAD 70 %
- AQ INSULIN SYRINGE 31G X 5/16" 1 ML
- ASSURE ID INSULIN SAFETY SYR 29G X 1/2" 1 ML
- ASSURE ID INSULIN SAFETY SYR 29G X 1/2" 0.5 ML (OTC)
- ASSURE ID INSULIN SAFETY SYR 31G X 15/64" 0.5 ML
- ASSURE ID INSULIN SAFETY SYR 31G X 15/64" 1 ML
- AUM ALCOHOL PREP PADS PAD 70 %
- BD ECLIPSE SYRINGE 30G X 1/2" 1 ML
- BD INSULIN SYR ULTRAFINE II 31G X 5/16" 0.3 ML
- BD INSULIN SYR ULTRAFINE II 31G X 5/16" 0.5 ML
- BD INSULIN SYR ULTRAFINE II 31G X 5/16" 1 ML
- BD INSULIN SYRINGE 27.5G X 5/8" 2 ML
- BD INSULIN SYRINGE 25G X 1" 1 ML
- BD INSULIN SYRINGE 25G X 5/8" 1 ML
- BD INSULIN SYRINGE 26G X 1/2" 1 ML
- BD INSULIN SYRINGE 27G X 1/2" 1 ML
- BD INSULIN SYRINGE 29G X 1/2" 0.5 ML (OTC)
- BD INSULIN SYRINGE 29G X 1/2" 0.5 ML (RX)
- BD INSULIN SYRINGE 29G X 1/2" 1 ML (OTC)
- BD INSULIN SYRINGE 29G X 1/2" 1 ML (RX)
- BD INSULIN SYRINGE HALF-UNIT 31G X 5/16" 0.3 ML
- BD INSULIN SYRINGE MICROFINE 27G X 5/8" 1 ML
- BD INSULIN SYRINGE MICROFINE 28G X 1/2" 0.5 ML
- BD INSULIN SYRINGE MICROFINE 28G X 1/2" 1 ML (OTC)
- BD INSULIN SYRINGE MICROFINE 28G X 1/2" 1 ML (RX)
- BD INSULIN SYRINGE U-100 1 ML
- BD INSULIN SYRINGE U-500
- BD INSULIN SYRINGE ULTRAFINE 29G X 1/2" 0.3 ML

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

- BD INSULIN SYRINGE ULTRAFINE 29G X 1/2" 0.5 ML
- BD INSULIN SYRINGE ULTRAFINE 29G X 1/2" 1 ML
- BD INSULIN SYRINGE ULTRAFINE 30G X 1/2" 0.3 ML
- BD INSULIN SYRINGE ULTRAFINE 30G X 1/2" 0.5 ML
- BD INSULIN SYRINGE ULTRAFINE 30G X 1/2" 1 ML
- BD INSULIN SYRINGE ULTRAFINE 31G X 5/16" 0.3 ML
- BD INSULIN SYRINGE ULTRAFINE 31G X 5/16" 0.5 ML
- BD SAFETY-LOK INSULIN SYRINGE 29G X 1/2" 1 ML
- BD SAFETYGLIDE INSULIN SYRINGE 29G X 1/2" 0.3 ML
- BD SAFETYGLIDE INSULIN SYRINGE 29G X 1/2" 0.5 ML
- BD SAFETYGLIDE INSULIN SYRINGE 30G X 5/16" 0.5 ML
- BD SAFETYGLIDE INSULIN SYRINGE 31G X 15/64" 0.3 ML
- BD SAFETYGLIDE INSULIN SYRINGE 31G X 15/64" 0.5 ML
- BD SAFETYGLIDE INSULIN SYRINGE 31G X 15/64" 1 ML
- BD SAFETYGLIDE INSULIN SYRINGE 31G X 5/16" 0.3 ML
- BD SAFETYGLIDE SYRINGE/NEEDLE 27G X 5/8" 1 ML
- BD SWAB SINGLE USE REGULAR PAD
- BD SWABS SINGLE USE BUTTERFLY PAD
- BD VEO INSULIN SYR U/F 1/2UNIT 31G X 15/64" 0.3 ML
- BD VEO INSULIN SYR ULTRAFINE 31G X 15/64" 0.3 ML
- BD VEO INSULIN SYR ULTRAFINE 31G X 15/64" 0.5 ML
- BD VEO INSULIN SYR ULTRAFINE 31G X 15/64" 1 ML
- BD VEO INSULIN SYRINGE U/F 31G X 15/64" 0.3 ML
- BD VEO INSULIN SYRINGE U/F 31G X 15/64" 0.5 ML
- BD VEO INSULIN SYRINGE U/F 31G X 15/64" 1 ML
- CAREONE INSULIN SYRINGE 30G X 1/2" 0.3 ML
- CAREONE INSULIN SYRINGE 30G X 1/2" 0.5 ML
- CAREONE INSULIN SYRINGE 30G X 1/2" 1 ML
- CAREONE INSULIN SYRINGE 31G X 5/16" 0.3 ML
- CAREONE INSULIN SYRINGE 31G X 5/16" 0.5 ML
- CAREONE INSULIN SYRINGE 31G X 5/16" 1 ML
- CARETOUCH ALCOHOL PREP PAD 70 %
- CARETOUCH INSULIN SYRINGE 28G X 5/16" 1 ML
- CARETOUCH INSULIN SYRINGE 29G X 5/16" 1 ML
- CARETOUCH INSULIN SYRINGE 30G X 5/16" 0.5 ML
- CARETOUCH INSULIN SYRINGE 30G X 5/16" 1 ML
- CARETOUCH INSULIN SYRINGE 31G X 5/16" 0.3 ML
- CARETOUCH INSULIN SYRINGE 31G X 5/16" 0.5 ML
- CARETOUCH INSULIN SYRINGE 31G X 5/16" 1 ML
- COMFORT ASSIST INSULIN SYRINGE 29G X 1/2" 1 ML
- COMFORT ASSIST INSULIN SYRINGE 31G X 5/16" 0.3 ML
- COMFORT EZ INSULIN SYRINGE 28G X 1/2" 0.5 ML
- COMFORT EZ INSULIN SYRINGE 28G X 1/2" 1 ML
- COMFORT EZ INSULIN SYRINGE 29G X 1/2" 0.3 ML

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

- COMFORT EZ INSULIN SYRINGE 29G X 1/2" 0.5 ML
- COMFORT EZ INSULIN SYRINGE 29G X 1/2" 1 ML
- COMFORT EZ INSULIN SYRINGE 30G X 1/2" 0.3 ML
- COMFORT EZ INSULIN SYRINGE 30G X 1/2" 0.5 ML
- COMFORT EZ INSULIN SYRINGE 30G X 1/2" 1 ML
- COMFORT EZ INSULIN SYRINGE 30G X 5/16" 0.3 ML
- COMFORT EZ INSULIN SYRINGE 30G X 5/16" 0.5 ML
- COMFORT EZ INSULIN SYRINGE 30G X 5/16" 1 ML
- COMFORT EZ INSULIN SYRINGE 31G X 15/64" 0.3 ML
- COMFORT EZ INSULIN SYRINGE 31G X 15/64" 0.5 ML
- COMFORT EZ INSULIN SYRINGE 31G X 15/64" 1 ML
- COMFORT EZ INSULIN SYRINGE 31G X 5/16" 0.3 ML
- COMFORT EZ INSULIN SYRINGE 31G X 5/16" 0.5 ML
- COMFORT EZ INSULIN SYRINGE 31G X 5/16" 1 ML
- COMFORT EZ INSULIN SYRINGE 31G X 5/16" 1 ML
- CURITY ALCOHOL PREPS PAD 70 %
- CURITY ALL PURPOSE SPONGES PAD 2"X2"
- CURITY GAUZE PAD 2"X2"
- CURITY GAUZE SPONGE PAD 2"X2"
- CURITY SPONGES PAD 2"X2"
- CVS GAUZE PAD 2"X2"
- CVS GAUZE STERILE PAD 2"X2"
- DERMACEA GAUZE SPONGE PAD 2"X2"
- DERMACEA IV DRAIN SPONGES PAD 2"X2"
- DERMACEA NON-WOVEN SPONGES PAD 2"X2"
- DERMACEA TYPE VII GAUZE PAD 2"X2"
- DROPLET INSULIN SYRINGE 29G X 1/2" 0.3 ML
- DROPLET INSULIN SYRINGE 29G X 1/2" 0.5 ML
- DROPLET INSULIN SYRINGE 29G X 1/2" 1 ML
- DROPLET INSULIN SYRINGE 30G X 1/2" 0.3 ML
- DROPLET INSULIN SYRINGE 30G X 1/2" 0.5 ML
- DROPLET INSULIN SYRINGE 30G X 1/2" 1 ML
- DROPLET INSULIN SYRINGE 30G X 15/64" 0.3 ML
- DROPLET INSULIN SYRINGE 30G X 15/64" 0.5 ML
- DROPLET INSULIN SYRINGE 30G X 15/64" 1 ML
- DROPLET INSULIN SYRINGE 30G X 5/16" 0.3 ML
- DROPLET INSULIN SYRINGE 30G X 5/16" 0.5 ML
- DROPLET INSULIN SYRINGE 30G X 5/16" 1 ML
- DROPLET INSULIN SYRINGE 31G X 1/4" 0.3 ML
- DROPLET INSULIN SYRINGE 31G X 1/4" 0.5 ML
- DROPLET INSULIN SYRINGE 31G X 1/4" 1 ML
- DROPLET INSULIN SYRINGE 31G X 15/64" 0.3 ML
- DROPLET INSULIN SYRINGE 31G X 15/64" 0.5 ML
- DROPLET INSULIN SYRINGE 31G X 15/64" 1 ML
- DROPLET INSULIN SYRINGE 31G X 5/16" 0.3 ML
- DROPLET INSULIN SYRINGE 31G X 5/16" 0.5 ML
- DROPLET INSULIN SYRINGE 31G X 5/16" 1 ML
- DROPSAFE ALCOHOL PREP PAD 70 %

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

- DROPSAFE SAFETY SYRINGE/NEEDLE 29G X 1/2" 1 ML
- DROPSAFE SAFETY SYRINGE/NEEDLE 31G X 15/64" 0.3 ML
- DROPSAFE SAFETY SYRINGE/NEEDLE 31G X 15/64" 0.5 ML
- DROPSAFE SAFETY SYRINGE/NEEDLE 31G X 15/64" 1 ML
- DROPSAFE SAFETY SYRINGE/NEEDLE 31G X 5/16" 0.3 ML
- DROPSAFE SAFETY SYRINGE/NEEDLE 31G X 5/16" 0.5 ML
- DROPSAFE SAFETY SYRINGE/NEEDLE 31G X 5/16" 1 ML
- DRUG MART ULTRA COMFORT SYR 29G X 1/2" 0.3 ML
- DRUG MART ULTRA COMFORT SYR 29G X 1/2" 1 ML
- DRUG MART ULTRA COMFORT SYR 30G X 5/16" 0.5 ML
- DRUG MART ULTRA COMFORT SYR 30G X 5/16" 1 ML
- EASY COMFORT ALCOHOL PADS PAD
- EASY COMFORT INSULIN SYRINGE 29G X 5/16" 0.5 ML
- EASY COMFORT INSULIN SYRINGE 29G X 5/16" 1 ML
- EASY COMFORT INSULIN SYRINGE 30G X 1/2" 0.5 ML
- EASY COMFORT INSULIN SYRINGE 30G X 1/2" 1 ML
- EASY COMFORT INSULIN SYRINGE 30G X 5/16" 0.5 ML
- EASY COMFORT INSULIN SYRINGE 30G X 5/16" 1 ML
- EASY COMFORT INSULIN SYRINGE 31G X 1/2" 0.3 ML
- EASY COMFORT INSULIN SYRINGE 31G X 5/16" 0.3 ML
- EASY COMFORT INSULIN SYRINGE 31G X 5/16" 0.5 ML
- EASY COMFORT INSULIN SYRINGE 31G X 5/16" 1 ML
- EASY COMFORT INSULIN SYRINGE 32G X 5/16" 0.5 ML
- EASY COMFORT INSULIN SYRINGE 32G X 5/16" 1 ML
- EASY TOUCH ALCOHOL PREP MEDIUM PAD 70 %
- EASY TOUCH FLIPLOCK INSULIN SY 29G X 1/2" 1 ML
- EASY TOUCH FLIPLOCK INSULIN SY 30G X 1/2" 1 ML
- EASY TOUCH FLIPLOCK INSULIN SY 30G X 5/16" 1 ML
- EASY TOUCH FLIPLOCK INSULIN SY 31G X 5/16" 1 ML
- EASY TOUCH FLIPLOCK SAFETY SYR 27G X 1/2" 1 ML
- EASY TOUCH INSULIN BARRELS U-100 1 ML
- EASY TOUCH INSULIN SAFETY SYR 29G X 1/2" 0.5 ML
- EASY TOUCH INSULIN SAFETY SYR 29G X 1/2" 1 ML
- EASY TOUCH INSULIN SAFETY SYR 30G X 1/2" 1 ML
- EASY TOUCH INSULIN SAFETY SYR 30G X 5/16" 0.5 ML
- EASY TOUCH INSULIN SYRINGE 27G X 1/2" 0.5 ML
- EASY TOUCH INSULIN SYRINGE 27G X 1/2" 1 ML
- EASY TOUCH INSULIN SYRINGE 27G X 5/8" 1 ML
- EASY TOUCH INSULIN SYRINGE 28G X 1/2" 0.5 ML
- EASY TOUCH INSULIN SYRINGE 28G X 1/2" 1 ML
- EASY TOUCH INSULIN SYRINGE 29G X 1/2" 0.5 ML
- EASY TOUCH INSULIN SYRINGE 29G X 1/2" 1 ML
- EASY TOUCH INSULIN SYRINGE 30G X 1/2" 0.3 ML

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

- EASY TOUCH INSULIN SYRINGE 30G X 1/2" 0.5 ML
- EASY TOUCH INSULIN SYRINGE 30G X 1/2" 1 ML
- EASY TOUCH INSULIN SYRINGE 30G X 5/16" 0.3 ML
- EASY TOUCH INSULIN SYRINGE 30G X 5/16" 0.5 ML
- EASY TOUCH INSULIN SYRINGE 30G X 5/16" 1 ML
- EASY TOUCH INSULIN SYRINGE 31G X 5/16" 0.3 ML
- EASY TOUCH INSULIN SYRINGE 31G X 5/16" 0.5 ML
- EASY TOUCH INSULIN SYRINGE 31G X 5/16" 1 ML
- EASY TOUCH SHEATHLOCK SYRINGE 29G X 1/2" 1 ML
- EASY TOUCH SHEATHLOCK SYRINGE 30G X 1/2" 1 ML
- EASY TOUCH SHEATHLOCK SYRINGE 30G X 5/16" 1 ML
- EASY TOUCH SHEATHLOCK SYRINGE 31G X 5/16" 1 ML
- EMBECTA INS SYR U/F 1/2 UNIT 31G X 15/64" 0.3 ML
- EMBECTA INS SYR U/F 1/2 UNIT 31G X 5/16" 0.3 ML
- EMBECTA INSULIN SYR ULTRAFINE 30G X 1/2" 0.3 ML
- EMBECTA INSULIN SYR ULTRAFINE 30G X 1/2" 0.5 ML
- EMBECTA INSULIN SYR ULTRAFINE 30G X 1/2" 1 ML
- EMBECTA INSULIN SYR ULTRAFINE 31G X 5/16" 0.5 ML
- EMBECTA INSULIN SYR ULTRAFINE 31G X 5/16" 1 ML
- EMBECTA INSULIN SYRINGE 28G X 1/2" 0.5 ML
- EMBECTA INSULIN SYRINGE U-100 27G X 5/8" 1 ML
- EMBECTA INSULIN SYRINGE U-100 28G X 1/2" 1 ML
- EQL ALCOHOL SWABS PAD 70 %
- EQL GAUZE PAD 2"X2"
- EQL INSULIN SYRINGE 30G X 5/16" 0.3 ML
- EQL INSULIN SYRINGE 30G X 5/16" 0.5 ML
- EQL INSULIN SYRINGE 30G X 5/16" 1 ML
- FREESTYLE PRECISION INS SYR 30G X 5/16" 0.5 ML
- FREESTYLE PRECISION INS SYR 30G X 5/16" 1 ML
- FREESTYLE PRECISION INS SYR 31G X 5/16" 0.5 ML
- FREESTYLE PRECISION INS SYR 31G X 5/16" 1 ML
- GAUZE PADS PAD 2"X2"
- GAUZE TYPE VII MEDI-PAK PAD 2"X2"
- GLOBAL ALCOHOL PREP EASE
- GLOBAL EASY GLIDE INSULIN SYR 31G X 15/64" 0.3 ML
- GLOBAL EASY GLIDE INSULIN SYR 31G X 15/64" 0.5 ML
- GLOBAL EASY GLIDE INSULIN SYR 31G X 15/64" 1 ML
- GLOBAL INJECT EASE INSULIN SYR 28G X 1/2" 0.5 ML
- GLOBAL INJECT EASE INSULIN SYR 28G X 1/2" 1 ML
- GLOBAL INJECT EASE INSULIN SYR 29G X 1/2" 0.5 ML
- GLOBAL INJECT EASE INSULIN SYR 29G X 1/2" 1 ML
- GLOBAL INJECT EASE INSULIN SYR 30G X 5/16" 0.5 ML
- GLOBAL INJECT EASE INSULIN SYR 30G X 5/16" 1 ML
- GLUCOPRO INSULIN SYRINGE 30G X 1/2" 0.3 ML
- GLUCOPRO INSULIN SYRINGE 30G X 1/2" 0.5 ML
- GLUCOPRO INSULIN SYRINGE 30G X 1/2" 1 ML

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

- GLUCOPRO INSULIN SYRINGE 30G X 5/16" 0.3 ML
- GLUCOPRO INSULIN SYRINGE 30G X 5/16" 0.5 ML
- GLUCOPRO INSULIN SYRINGE 30G X 5/16" 1 ML
- GLUCOPRO INSULIN SYRINGE 31G X 5/16" 0.3 ML
- GLUCOPRO INSULIN SYRINGE 31G X 5/16" 0.5 ML
- GLUCOPRO INSULIN SYRINGE 31G X 5/16" 1 ML
- GNP ALCOHOL SWABS PAD
- GNP INSULIN SYRINGE 28G X 1/2" 1 ML
- GNP INSULIN SYRINGE 29G X 1/2" 0.5 ML
- GNP INSULIN SYRINGE 29G X 1/2" 1 ML
- GNP INSULIN SYRINGE 30G X 5/16" 0.5 ML
- GNP INSULIN SYRINGE 30G X 5/16" 1 ML
- GNP INSULIN SYRINGES 29GX1/2" 29G X 1/2" 0.5 ML
- GNP INSULIN SYRINGES 29GX1/2" 29G X 1/2" 1 ML
- GNP INSULIN SYRINGES 30G X 5/16" 1 ML
- GNP INSULIN SYRINGES 30GX5/16" 30G X 5/16" 0.3 ML
- GNP INSULIN SYRINGES 31GX5/16" 31G X 5/16" 0.3 ML
- GNP STERILE GAUZE PAD 2"X2"
- GNP ULTRA COM INSULIN SYRINGE 29G X 1/2" 0.3 ML
- GNP ULTRA COM INSULIN SYRINGE 30G X 5/16" 0.3 ML
- GOODSENSE ALCOHOL SWABS PAD 70 %
- H-E-B INCONTROL ALCOHOL PAD
- HEALTHWISE INSULIN SYR/NEEDLE 30G X 5/16" 0.3 ML
- HEALTHWISE INSULIN SYR/NEEDLE 30G X 5/16" 0.5 ML
- HEALTHWISE INSULIN SYR/NEEDLE 30G X 5/16" 1 ML
- HEALTHWISE INSULIN SYR/NEEDLE 31G X 5/16" 0.3 ML
- HEALTHWISE INSULIN SYR/NEEDLE 31G X 5/16" 0.5 ML
- HEALTHWISE INSULIN SYR/NEEDLE 31G X 5/16" 1 ML
- HM STERILE PADS PAD 2"X2"
- HM ULTICARE INSULIN SYRINGE 30G X 1/2" 1 ML
- HM ULTICARE INSULIN SYRINGE 31G X 5/16" 0.3 ML
- INSULIN SYRINGE 29G X 1/2" 1 ML
- INSULIN SYRINGE 30G X 5/16" 1 ML
- INSULIN SYRINGE 31G X 5/16" 0.3 ML
- INSULIN SYRINGE 31G X 5/16" 0.5 ML
- INSULIN SYRINGE-NEEDLE U-100 27G X 1/2" 0.5 ML (OTC)
- INSULIN SYRINGE-NEEDLE U-100 27G X 1/2" 0.5 ML (RX)
- INSULIN SYRINGE-NEEDLE U-100 27G X 1/2" 1 ML (RX)
- INSULIN SYRINGE-NEEDLE U-100 28G X 1/2" 0.5 ML (RX)
- INSULIN SYRINGE-NEEDLE U-100 28G X 1/2" 1 ML (RX)
- INSULIN SYRINGE-NEEDLE U-100 30G X 5/16" 1 ML
- INSULIN SYRINGE-NEEDLE U-100 31G X 1/4" 0.3 ML
- INSULIN SYRINGE-NEEDLE U-100 31G X 1/4" 0.5 ML
- INSULIN SYRINGE-NEEDLE U-100 31G X 1/4" 1 ML
- INSULIN SYRINGE-NEEDLE U-100 31G X 5/16" 0.5 ML (OTC)
- INSULIN SYRINGE/NEEDLE 27G X 1/2" 0.5 ML

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

- INSULIN SYRINGE/NEEDLE 28G X 1/2" 0.5 ML
- INSULIN SYRINGE/NEEDLE 28G X 1/2" 1 ML
- J & J GAUZE PAD 2"X2"
- KENDALL HYDROPHILIC FOAM DRESS PAD 2"X2"
- KENDALL HYDROPHILIC FOAM PLUS PAD 2"X2"
- KINRAY INSULIN SYRINGE 29G X 1/2" 0.5 ML
- KMART VALU INSULIN SYRINGE 29G U-100 1 ML
- KMART VALU INSULIN SYRINGE 30G U-100 0.3 ML
- KMART VALU INSULIN SYRINGE 30G U-100 1 ML
- LITETOUCH INSULIN SYRINGE 28G X 1/2" 0.5 ML
- LITETOUCH INSULIN SYRINGE 28G X 1/2" 1 ML
- LITETOUCH INSULIN SYRINGE 29G X 1/2" 0.3 ML
- LITETOUCH INSULIN SYRINGE 29G X 1/2" 0.5 ML
- LITETOUCH INSULIN SYRINGE 29G X 1/2" 1 ML
- LITETOUCH INSULIN SYRINGE 30G X 5/16" 0.3 ML
- LITETOUCH INSULIN SYRINGE 30G X 5/16" 0.5 ML
- LITETOUCH INSULIN SYRINGE 30G X 5/16" 1 ML
- LITETOUCH INSULIN SYRINGE 31G X 5/16" 0.3 ML
- LITETOUCH INSULIN SYRINGE 31G X 5/16" 0.5 ML
- LITETOUCH INSULIN SYRINGE 31G X 5/16" 1 ML
- MAGELLAN INSULIN SAFETY SYR 29G X 1/2" 0.3 ML
- MAGELLAN INSULIN SAFETY SYR 29G X 1/2" 0.5 ML
- MAGELLAN INSULIN SAFETY SYR 29G X 1/2" 1 ML
- MAGELLAN INSULIN SAFETY SYR 30G X 5/16" 0.3 ML
- MAGELLAN INSULIN SAFETY SYR 30G X 5/16" 0.5 ML
- MAGELLAN INSULIN SAFETY SYR 30G X 5/16" 1 ML
- MAXI-COMFORT INSULIN SYRINGE 28G X 1/2" 0.5 ML
- MAXI-COMFORT INSULIN SYRINGE 28G X 1/2" 1 ML
- MAXICOMFORT SYR 27G X 1/2" 27G X 1/2" 0.5 ML
- MAXICOMFORT SYR 27G X 1/2" 27G X 1/2" 1 ML
- MEDIC INSULIN SYRINGE 30G X 5/16" 0.3 ML
- MEDIC INSULIN SYRINGE 30G X 5/16" 0.5 ML
- MEDPURA ALCOHOL PADS 70 % EXTERNAL
- MEIJER ALCOHOL SWABS PAD 70 %
- MIRASORB SPONGES 2"X2"
- MONOJECT INSULIN SYRINGE 25G X 5/8" 1 ML
- MONOJECT INSULIN SYRINGE 27G X 1/2" 1 ML (OTC)
- MONOJECT INSULIN SYRINGE 28G X 1/2" 0.5 ML (RX)
- MONOJECT INSULIN SYRINGE 28G X 1/2" 1 ML (OTC)
- MONOJECT INSULIN SYRINGE 28G X 1/2" 1 ML (RX)
- MONOJECT INSULIN SYRINGE 29G X 1/2" 0.3 ML
- MONOJECT INSULIN SYRINGE 29G X 1/2" 0.5 ML
- MONOJECT INSULIN SYRINGE 29G X 1/2" 1 ML (RX)
- MONOJECT INSULIN SYRINGE 30G X 5/16" 0.3 ML
- MONOJECT INSULIN SYRINGE 30G X 5/16" 0.5 ML (RX)

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

- MONOJECT INSULIN SYRINGE 30G X 5/16" 1 ML (RX)
- MONOJECT INSULIN SYRINGE 31G X 5/16" 1 ML
- MONOJECT INSULIN SYRINGE U-100 1 ML
- MONOJECT ULTRA COMFORT SYRINGE 28G X 1/2" 0.5 ML (OTC)
- MONOJECT ULTRA COMFORT SYRINGE 28G X 1/2" 0.5 ML (RX)
- MONOJECT ULTRA COMFORT SYRINGE 28G X 1/2" 1 ML (OTC)
- MONOJECT ULTRA COMFORT SYRINGE 29G X 1/2" 0.5 ML
- MONOJECT ULTRA COMFORT SYRINGE 29G X 1/2" 1 ML
- MONOJECT ULTRA COMFORT SYRINGE 30G X 5/16" 0.3 ML (OTC)
- MONOJECT ULTRA COMFORT SYRINGE 30G X 5/16" 0.3 ML (RX)
- MONOJECT ULTRA COMFORT SYRINGE 30G X 5/16" 0.5 ML (RX)
- PRECISION SURE-DOSE SYRINGE 28G X 1/2" 0.5 ML
- PRECISION SURE-DOSE SYRINGE 28G X 1/2" 1 ML
- PRECISION SURE-DOSE SYRINGE 29G X 1/2" 0.5 ML
- PRECISION SURE-DOSE SYRINGE 30G X 3/8" 0.5 ML
- PRECISION SURE-DOSE SYRINGE 30G X 5/16" 0.3 ML
- PRECISION SUREDOSE PLUS SYR 29G X 1/2" 0.3 ML
- PRECISION SUREDOSE PLUS SYR 29G X 1/2" 1 ML
- PREFERRED PLUS INSULIN SYRINGE 28G X 1/2" 0.5 ML
- PRO COMFORT ALCOHOL PAD 70 %
- PRO COMFORT INSULIN SYRINGE 30G X 1/2" 0.5 ML
- PRO COMFORT INSULIN SYRINGE 30G X 1/2" 1 ML
- PRO COMFORT INSULIN SYRINGE 30G X 5/16" 0.5 ML
- PRO COMFORT INSULIN SYRINGE 30G X 5/16" 1 ML
- PRO COMFORT INSULIN SYRINGE 31G X 5/16" 0.5 ML
- PRO COMFORT INSULIN SYRINGE 31G X 5/16" 1 ML
- PRODIGY INSULIN SYRINGE 28G X 1/2" 1 ML
- PRODIGY INSULIN SYRINGE 31G X 5/16" 0.3 ML
- PRODIGY INSULIN SYRINGE 31G X 5/16" 0.5 ML
- PURE COMFORT ALCOHOL PREP PAD
- QC ALCOHOL
- QC ALCOHOL SWABS PAD 70 %
- QC BORDER ISLAND GAUZE PAD 2"X2"
- RA ALCOHOL SWABS PAD 70 %
- RA INSULIN SYRINGE 29G X 1/2" 1 ML
- RA INSULIN SYRINGE 30G X 5/16" 0.5 ML
- RA INSULIN SYRINGE 30G X 5/16" 1 ML
- *ra isopropyl alcohol wipes*
- RA STERILE PAD 2"X2"
- REALITY INSULIN SYRINGE 28G X 1/2" 0.5 ML
- REALITY INSULIN SYRINGE 28G X 1/2" 1 ML
- REALITY INSULIN SYRINGE 29G X 1/2" 0.5 ML
- REALITY INSULIN SYRINGE 29G X 1/2" 1 ML
- REALITY SWABS PAD
- RELI-ON INSULIN SYRINGE 29G 0.3 ML
- RELI-ON INSULIN SYRINGE 29G 0.5 ML
- RELI-ON INSULIN SYRINGE 29G X 1/2" 1 ML

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

- RELION ALCOHOL SWABS PAD
- RELION INSULIN SYRINGE 31G X 15/64" 0.3 ML
- RELION INSULIN SYRINGE 31G X 15/64" 0.5 ML
- RELION INSULIN SYRINGE 31G X 15/64" 1 ML
- RESTORE CONTACT LAYER PAD 2"X2"
- SAFETY INSULIN SYRINGES 29G X 1/2" 0.5 ML
- SAFETY INSULIN SYRINGES 29G X 1/2" 1 ML
- SAFETY INSULIN SYRINGES 30G X 1/2" 1 ML
- SAFETY INSULIN SYRINGES 30G X 5/16" 0.5 ML
- SB ALCOHOL PREP PAD 70 %
- SB INSULIN SYRINGE 29G X 1/2" 0.5 ML
- SB INSULIN SYRINGE 29G X 1/2" 1 ML
- SB INSULIN SYRINGE 30G X 5/16" 0.5 ML
- SB INSULIN SYRINGE 30G X 5/16" 1 ML
- SB INSULIN SYRINGE 31G X 5/16" 1 ML
- SECURESAFE INSULIN SYRINGE 29G X 1/2" 0.5 ML
- SECURESAFE INSULIN SYRINGE 29G X 1/2" 1 ML
- SM ALCOHOL PREP PAD
- SM ALCOHOL PREP PAD 6-70 % EXTERNAL
- SM ALCOHOL PREP PAD 70 %
- SM GAUZE PAD 2"X2"
- STERILE GAUZE PAD 2"X2"
- STERILE PAD 2"X2"
- SURE COMFORT ALCOHOL PREP PAD 70 %
- SURE COMFORT INSULIN SYRINGE 28G X 1/2" 0.5 ML
- SURE COMFORT INSULIN SYRINGE 28G X 1/2" 1 ML
- SURE COMFORT INSULIN SYRINGE 29G X 1/2" 0.3 ML
- SURE COMFORT INSULIN SYRINGE 29G X 1/2" 0.5 ML
- SURE COMFORT INSULIN SYRINGE 29G X 1/2" 1 ML
- SURE COMFORT INSULIN SYRINGE 30G X 1/2" 0.3 ML
- SURE COMFORT INSULIN SYRINGE 30G X 1/2" 0.5 ML
- SURE COMFORT INSULIN SYRINGE 30G X 1/2" 1 ML
- SURE COMFORT INSULIN SYRINGE 30G X 5/16" 0.3 ML
- SURE COMFORT INSULIN SYRINGE 30G X 5/16" 0.5 ML
- SURE COMFORT INSULIN SYRINGE 30G X 5/16" 1 ML
- SURE COMFORT INSULIN SYRINGE 31G X 1/4" 0.3 ML
- SURE COMFORT INSULIN SYRINGE 31G X 1/4" 0.5 ML
- SURE COMFORT INSULIN SYRINGE 31G X 1/4" 1 ML
- SURE COMFORT INSULIN SYRINGE 31G X 5/16" 0.3 ML
- SURE COMFORT INSULIN SYRINGE 31G X 5/16" 0.5 ML
- SURE COMFORT INSULIN SYRINGE 31G X 5/16" 1 ML
- SURE-JECT INSULIN SYRINGE 31G X 5/16" 0.3 ML
- SURE-JECT INSULIN SYRINGE 31G X 5/16" 0.5 ML
- SURE-JECT INSULIN SYRINGE 31G X 5/16" 1 ML
- SURE-PREP ALCOHOL PREP PAD 70 %
- SURGICAL GAUZE SPONGE PAD 2"X2"
- TECHLITE INSULIN SYRINGE 29G X 1/2" 0.5 ML

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

- TERUMO INSULIN SYRINGE 29G X 1/2" 0.3 ML
- THERAGAUZE PAD 2"X2"
- TOPCARE ULTRA COMFORT INS SYR 29G X 1/2" 0.3 ML
- TOPCARE ULTRA COMFORT INS SYR 29G X 1/2" 0.5 ML
- TOPCARE ULTRA COMFORT INS SYR 29G X 1/2" 1 ML
- TOPCARE ULTRA COMFORT INS SYR 30G X 5/16" 0.3 ML
- TOPCARE ULTRA COMFORT INS SYR 30G X 5/16" 0.5 ML
- TOPCARE ULTRA COMFORT INS SYR 30G X 5/16" 1 ML
- TOPCARE ULTRA COMFORT INS SYR 31G X 5/16" 0.3 ML
- TOPCARE ULTRA COMFORT INS SYR 31G X 5/16" 0.5 ML
- TOPCARE ULTRA COMFORT INS SYR 31G X 5/16" 1 ML
- TRUE COMFORT ALCOHOL PREP PADS PAD 70 %
- TRUE COMFORT INSULIN SYRINGE 30G X 1/2" 0.5 ML
- TRUE COMFORT INSULIN SYRINGE 30G X 1/2" 1 ML
- TRUE COMFORT INSULIN SYRINGE 30G X 5/16" 0.5 ML
- TRUE COMFORT INSULIN SYRINGE 30G X 5/16" 1 ML
- TRUE COMFORT INSULIN SYRINGE 31G X 5/16" 0.5 ML
- TRUE COMFORT INSULIN SYRINGE 31G X 5/16" 1 ML
- TRUE COMFORT INSULIN SYRINGE 32G X 5/16" 1 ML
- TRUE COMFORT PRO ALCOHOL PREP PAD 70 %
- TRUE COMFORT PRO INSULIN SYR 30G X 1/2" 0.5 ML
- TRUE COMFORT PRO INSULIN SYR 30G X 1/2" 1 ML
- TRUE COMFORT PRO INSULIN SYR 30G X 5/16" 0.5 ML
- TRUE COMFORT PRO INSULIN SYR 30G X 5/16" 1 ML
- TRUE COMFORT PRO INSULIN SYR 31G X 5/16" 0.5 ML
- TRUE COMFORT PRO INSULIN SYR 31G X 5/16" 1 ML
- TRUE COMFORT PRO INSULIN SYR 32G X 5/16" 0.5 ML
- TRUE COMFORT PRO INSULIN SYR 32G X 5/16" 1 ML
- TRUEPLUS INSULIN SYRINGE 28G X 1/2" 0.5 ML
- TRUEPLUS INSULIN SYRINGE 28G X 1/2" 1 ML
- TRUEPLUS INSULIN SYRINGE 29G X 1/2" 0.3 ML
- TRUEPLUS INSULIN SYRINGE 29G X 1/2" 0.5 ML
- TRUEPLUS INSULIN SYRINGE 29G X 1/2" 1 ML
- TRUEPLUS INSULIN SYRINGE 30G X 5/16" 0.3 ML
- TRUEPLUS INSULIN SYRINGE 30G X 5/16" 0.5 ML
- TRUEPLUS INSULIN SYRINGE 30G X 5/16" 1 ML
- TRUEPLUS INSULIN SYRINGE 31G X 5/16" 0.3 ML
- TRUEPLUS INSULIN SYRINGE 31G X 5/16" 0.5 ML
- TRUEPLUS INSULIN SYRINGE 31G X 5/16" 1 ML
- ULTICARE INSULIN SAFETY SYR 29G X 1/2" 0.5 ML
- ULTICARE INSULIN SAFETY SYR 29G X 1/2" 1 ML
- ULTICARE INSULIN SYRINGE 28G X 1/2" 0.5 ML
- ULTICARE INSULIN SYRINGE 28G X 1/2" 1 ML
- ULTICARE INSULIN SYRINGE 29G X 1/2" 0.3 ML

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C



- ULTRA FLO INSULIN SYR 1/2 UNIT 30G X 1/2" 0.3 ML
- ULTRA FLO INSULIN SYR 1/2 UNIT 30G X 5/16" 0.3 ML
- ULTRA FLO INSULIN SYR 1/2 UNIT 31G X 5/16" 0.3 ML
- ULTRA FLO INSULIN SYRINGE 29G X 1/2" 0.3 ML
- ULTRA FLO INSULIN SYRINGE 29G X 1/2" 0.5 ML
- ULTRA FLO INSULIN SYRINGE 29G X 1/2" 1 ML
- ULTRA FLO INSULIN SYRINGE 30G X 1/2" 0.3 ML
- ULTRA FLO INSULIN SYRINGE 30G X 1/2" 0.5 ML
- ULTRA FLO INSULIN SYRINGE 30G X 1/2" 1 ML
- ULTRA FLO INSULIN SYRINGE 30G X 5/16" 0.3 ML
- ULTRA FLO INSULIN SYRINGE 30G X 5/16" 0.5 ML
- ULTRA FLO INSULIN SYRINGE 30G X 5/16" 1 ML
- ULTRA FLO INSULIN SYRINGE 31G X 5/16" 0.3 ML
- ULTRA FLO INSULIN SYRINGE 31G X 5/16" 0.5 ML
- ULTRA FLO INSULIN SYRINGE 31G X 5/16" 1 ML
- ULTRA-COMFORT INSULIN SYRINGE 29G X 1/2" 0.5 ML
- ULTRA-THIN II INS SYR SHORT 30G X 5/16" 0.3 ML
- ULTRA-THIN II INS SYR SHORT 30G X 5/16" 0.5 ML
- ULTRA-THIN II INS SYR SHORT 30G X 5/16" 1 ML
- ULTRA-THIN II INS SYR SHORT 31G X 5/16" 0.3 ML
- ULTRA-THIN II INS SYR SHORT 31G X 5/16" 0.5 ML
- ULTRA-THIN II INS SYR SHORT 31G X 5/16" 1 ML
- ULTRA-THIN II INSULIN SYRINGE 29G X 1/2" 0.5 ML
- ULTRA-THIN II INSULIN SYRINGE 29G X 1/2" 1 ML
- ULTRACARE INSULIN SYRINGE 30G X 1/2" 0.5 ML
- ULTRACARE INSULIN SYRINGE 30G X 1/2" 1 ML
- ULTRACARE INSULIN SYRINGE 30G X 5/16" 0.3 ML
- ULTRACARE INSULIN SYRINGE 30G X 5/16" 0.5 ML
- ULTRACARE INSULIN SYRINGE 30G X 5/16" 1 ML
- ULTRACARE INSULIN SYRINGE 31G X 5/16" 0.3 ML
- ULTRACARE INSULIN SYRINGE 31G X 5/16" 0.5 ML
- ULTRACARE INSULIN SYRINGE 31G X 5/16" 1 ML
- VALUE HEALTH INSULIN SYRINGE 29G X 1/2" 0.5 ML
- VALUE HEALTH INSULIN SYRINGE 29G X 1/2" 1 ML
- VANISHPOINT INSULIN SYRINGE 29G X 5/16" 1 ML
- VANISHPOINT INSULIN SYRINGE 30G X 3/16" 0.5 ML
- VANISHPOINT INSULIN SYRINGE 30G X 3/16" 1 ML
- VANISHPOINT INSULIN SYRINGE 30G X 5/16" 0.5 ML
- VANISHPOINT INSULIN SYRINGE 30G X 5/16" 1 ML
- VERIFINE INSULIN SYRINGE 29G X 1/2" 0.5 ML
- VERIFINE INSULIN SYRINGE 29G X 1/2" 1 ML
- VERIFINE INSULIN SYRINGE 31G X 5/16" 0.3 ML
- VERIFINE INSULIN SYRINGE 31G X 5/16" 0.5 ML
- VERIFINE INSULIN SYRINGE 31G X 5/16" 1 ML

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

- VP INSULIN SYRINGE 29G X 1/2" 0.3 ML
- WEBCOL ALCOHOL PREP LARGE PAD 70 %
- ZEV RX STERILE ALCOHOL PREP PAD PAD 70 %

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	LIFETIME
<b>Other Criteria</b>	ONLY COVERED UNDER PART D WHEN USED CONCURRENTLY WITH INSULIN.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# INTERFERON FOR MS-AVONEX

## Products Affected

- AVONEX PEN INTRAMUSCULAR  
AUTO-INJECTOR KIT
- AVONEX PREFILLED  
INTRAMUSCULAR PREFILLED  
SYRINGE KIT

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# INTERFERON FOR MS-BETASERON

---

## Products Affected

- BETASERON SUBCUTANEOUS KIT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# INTERFERON FOR MS-PLEGRIDY

## Products Affected

- PLEGRIDY STARTER PACK SUBCUTANEOUS SOLUTION AUTO-INJECTOR
- PLEGRIDY STARTER PACK SUBCUTANEOUS SOLUTION PREFILLED SYRINGE
- PLEGRIDY SUBCUTANEOUS SOLUTION AUTO-INJECTOR
- PLEGRIDY SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# INTERFERON GAMMA-1B

## Products Affected

- ACTIMMUNE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: CHRONIC GRANULOMATOUS DISEASE (CGD): PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST, INFECTIOUS DISEASE SPECIALIST, OR IMMUNOLOGIST. SEVERE MALIGNANT OSTEOPETROSIS (SMO): PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST OR HEMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	RENEWAL: CGD, SMO: 1) DEMONSTRATED CLINICAL BENEFIT COMPARED TO BASELINE, AND 2) HAS NOT RECEIVED HEMATOPOIETIC CELL TRANSPLANTATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# IPILIMUMAB

## Products Affected

- YERVOY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: UNRESECT/MET MEL: 4MO, RCC/CRC/HCC: 3MO, ALL OTHERS: 12MO. INITIAL/RENEWAL: CUTAN MEL: 6MO
Other Criteria	RENEWAL: ADJUVANT CUTANEOUS MELANOMA: NO EVIDENCE OF DISEASE RECURRENCE (DEFINED AS THE APPEARANCE OF ONE OR MORE NEW MELANOMA LESIONS: LOCAL, REGIONAL OR DISTANT METASTASIS). THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# IVACAFTOR

## Products Affected

- KALYDECO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	CYSTIC FIBROSIS (CF): INITIAL: CONFIRMED MUTATION IN CFTR GENE ACCEPTABLE FOR THE TREATMENT OF CYSTIC FIBROSIS
Age Restrictions	
Prescriber Restrictions	CF: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR CYSTIC FIBROSIS EXPERT
Coverage Duration	INITIAL: 12 MONTHS. RENEWAL: LIFETIME
Other Criteria	CF: INITIAL: NOT HOMOZYGOUS FOR F508DEL MUTATION IN CFTR GENE. RENEWAL: IMPROVEMENT IN CLINICAL STATUS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# IVOSIDENIB

---

## Products Affected

- TIBSOVO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# IXAZOMIB

---

## Products Affected

- NINLARO

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# LANREOTIDE

## Products Affected

- LANREOTIDE ACETATE
- SOMATULINE DEPOT  
SUBCUTANEOUS SOLUTION 60  
MG/0.2ML, 90 MG/0.3ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	ACROMEGALY: INITIAL: THERAPY IS PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST.
Coverage Duration	ACROMEGALY: INITIAL: 3 MOS, RENEWAL: 12 MOS.GEP-NETS, CARCINOID SYNDROME: 12 MOS.
Other Criteria	ACROMEGALY: INITIAL: TRIAL OF OR CONTRAINDICATION TO ONE GENERIC OCTREOTIDE INJECTION. RENEWAL: 1) REDUCTION, NORMALIZATION, OR MAINTENANCE OF IGF-1 LEVELS BASED ON AGE AND GENDER, AND 2) IMPROVEMENT OR SUSTAINED REMISSION OF CLINICAL SYMPTOMS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# LAPATINIB

---

## Products Affected

- *lapatinib ditosylate*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# LAROTRECTINIB

## Products Affected

- VITRAKVI ORAL CAPSULE 100 MG, 25 MG
- VITRAKVI ORAL SOLUTION

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	VITRAKVI ORAL SOLUTION: 1) TRIAL OF VITRAKVI CAPSULES, OR 2) UNABLE TO TAKE CAPSULE FORMULATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# LAZERTINIB

## Products Affected

- LAZCLUZE ORAL TABLET 240 MG, 80 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# LEDIPASVIR-SOFOSBUVIR

## Products Affected

- HARVONI ORAL PACKET 33.75-150 MG, 45-200 MG
- HARVONI ORAL TABLET

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	HCV RNA LEVEL WITHIN PAST 6 MONTHS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE.
<b>Other Criteria</b>	1) CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE, AND 2) NOT CONCURRENTLY TAKING ANY OF THE FOLLOWING: CARBAMAZEPINE, PHENYTOIN, PHENOBARBITAL, OXCARBAZEPINE, RIFAMPIN, RIFABUTIN, RIFAPENTINE, ROSUVASTATIN, TIPRANAVIR/RITONAVIR, SOFOSBUVIR (AS A SINGLE AGENT), EPCLUSA, ZEPATIER, MAVYRET, OR VOSEVI. REQUESTS FOR HARVONI 45MG-200MG PELLETS: PATIENT IS UNABLE TO SWALLOW TABLETS.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# LENALIDOMIDE

---

## Products Affected

- *lenalidomide*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# LENVATINIB

## Products Affected

- LENVIMA (10 MG DAILY DOSE)
- LENVIMA (12 MG DAILY DOSE)
- LENVIMA (14 MG DAILY DOSE)
- LENVIMA (18 MG DAILY DOSE)
- LENVIMA (20 MG DAILY DOSE)
- LENVIMA (24 MG DAILY DOSE)
- LENVIMA (4 MG DAILY DOSE)
- LENVIMA (8 MG DAILY DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# LETERMIVIR

## Products Affected

- PREVYMIS ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	HSCT: NOT AT RISK FOR LATE CMV: 4 MOS, AT RISK FOR LATE CMV: 7 MOS. KIDNEY TRANSPLANT: 7 MOS.
Other Criteria	HEMATOPOIETIC STEM CELL TRANSPLANT (HSCT): 1) THERAPY WILL BE INITIATED BETWEEN DAY 0 AND DAY 28 POST TRANSPLANT, AND 2) WILL NOT RECEIVE THE MEDICATION BEYOND 100 DAYS POST TRANSPLANT IF NOT AT RISK FOR LATE CYTOMEGALOVIRUS (CMV) INFECTION AND DISEASE, OR BEYOND 200 DAYS POST TRANSPLANT IF AT RISK FOR LATE CMV INFECTION AND DISEASE. KIDNEY TRANSPLANT: 1) THERAPY WILL BE INITIATED BETWEEN DAY 0 AND DAY 7 POST TRANSPLANT, AND 2) WILL NOT RECEIVE THE MEDICATION BEYOND 200 DAYS POST TRANSPLANT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# LEUPROLIDE

---

## Products Affected

- *leuprolide acetate injection*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	PROSTATE CANCER: 12 MONTHS.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# LEUPROLIDE DEPOT

## Products Affected

- LEUPROLIDE ACETATE (3 MONTH)
- LUTRATE DEPOT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# LEUPROLIDE-ELIGARD

---

## Products Affected

- ELIGARD

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# LEUPROLIDE-LUPRON DEPOT

## Products Affected

- LUPRON DEPOT (1-MONTH)
- LUPRON DEPOT (3-MONTH)
- LUPRON DEPOT (4-MONTH)
- LUPRON DEPOT (6-MONTH)

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: ENDOMETRIOSIS: DIAGNOSIS IS CONFIRMED VIA SURGICAL OR DIRECT VISUALIZATION (E.G., PELVIC ULTRASOUND) OR HISTOPATHOLOGICAL CONFIRMATION (E.G., LAPAROSCOPY OR LAPAROTOMY) IN THE LAST 10 YEARS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: ENDOMETRIOSIS: PRESCRIBED BY OR IN CONSULTATION WITH AN OBSTETRICIAN/GYNECOLOGIST.
<b>Coverage Duration</b>	PROSTATE CA: 12 MOS. UTERINE FIBROIDS: 3 MOS. ENDOMETRIOSIS: INITIAL/RENEWAL: 6 MOS.
<b>Other Criteria</b>	INITIAL: ENDOMETRIOSIS: 1) NO CONCURRENT USE WITH ANOTHER GNRH-MODULATING AGENT, 2) TRIAL OF OR CONTRAINDICATION TO NSAID AND PROGESTIN-CONTAINING PREPARATION, AND 3) HAS NOT RECEIVED A TOTAL OF 12 MONTHS OF TREATMENT PER LIFETIME. RENEWAL: ENDOMETRIOSIS: 1) IMPROVEMENT OF PAIN RELATED TO ENDOMETRIOSIS WHILE ON THERAPY, 2) RECEIVING CONCOMITANT ADD-BACK THERAPY (I.E., COMBINATION ESTROGEN-PROGESTIN OR PROGESTIN-ONLY CONTRACEPTIVE PREPARATION), 3) NO CONCURRENT USE WITH ANOTHER GNRH-MODULATING AGENT, AND 4) HAS NOT RECEIVED A TOTAL OF 12 MONTHS OF TREATMENT PER LIFETIME. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# LEUPROLIDE-LUPRON DEPOT-PED

## Products Affected

- LUPRON DEPOT-PED (3-MONTH)
- LUPRON DEPOT-PED (6-MONTH)

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	CENTRAL PRECOCIOUS PUBERTY (CPP): INITIAL: FEMALES: ELEVATED LEVELS OF FOLLICLE-STIMULATING HORMONE (FSH) GREATER THAN 4.0 MIU/ML AND LUTEINIZING HORMONE (LH) LEVEL GREATER THAN 0.2 TO 0.3 MIU/ML AT DIAGNOSIS. MALES: ELEVATED LEVELS OF FSH GREATER THAN 5.0 MIU/ML AND LH LEVEL GREATER THAN 0.2 TO 0.3 MIU/ML AT DIAGNOSIS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	CPP: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	CPP: INITIAL: FEMALES: 1) YOUNGER THAN 8 YEARS OF AGE AT ONSET OF CPP, AND 2) AT TANNER STAGE 2 OR ABOVE FOR BREAST DEVELOPMENT AND PUBIC HAIR GROWTH. MALES: 1) YOUNGER THAN 9 YEARS OF AGE AT ONSET OF CPP, AND 2) AT TANNER STAGE 2 OR ABOVE FOR GENITAL DEVELOPMENT AND PUBIC HAIR GROWTH. RENEWAL: 1) TANNER STAGING AT INITIAL DIAGNOSIS HAS STABILIZED OR REGRESSED DURING THREE SEPARATE MEDICAL VISITS IN THE PREVIOUS YEAR, AND 2) HAS NOT REACHED ACTUAL AGE WHICH CORRESPONDS TO CURRENT PUBERTAL AGE. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# L-GLUTAMINE

## Products Affected

- *l-glutamine oral packet*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	SICKLE CELL DISEASE(SCD): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST
Coverage Duration	INITIAL: 12 MONTHS. RENEWAL: LIFETIME.
Other Criteria	SCD: INITIAL: AGES 18 YEARS OR OLDER: 1) AT LEAST 2 SICKLE CELL CRISES IN THE PAST YEAR, 2) SICKLE-CELL ASSOCIATED SYMPTOMS WHICH ARE INTERFERING WITH ACTIVITIES OF DAILY LIVING, OR 3) HISTORY OF OR HAS RECURRENT ACUTE CHEST SYNDROME. AGES 5 TO 17 YEARS: APPROVED WITHOUT ADDITIONAL CRITERIA. RENEWAL: MAINTAINED OR EXPERIENCED A REDUCTION IN ACUTE COMPLICATIONS OF SCD.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# LONCASTUXIMAB TESIRINE-LPYL

---

## Products Affected

- ZYNLONTA

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# LORLATINIB

## Products Affected

- LORBRENA ORAL TABLET 100 MG,  
25 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# LOTILANER

---

## Products Affected

- XDEMVY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	DEMODEX BLEPHARITIS: 18 YEARS OF AGE OR OLDER
Prescriber Restrictions	
Coverage Duration	6 WEEKS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# LUMACAFITOR-IVACAFITOR

## Products Affected

- ORKAMBI ORAL TABLET

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: CYSTIC FIBROSIS (CF): CONFIRMED MUTATION IN CFTR GENE ACCEPTABLE FOR THE TREATMENT OF CF.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	CF: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR CF EXPERT.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS, RENEWAL: LIFETIME.
<b>Other Criteria</b>	CF: RENEWAL: IMPROVEMENT IN CLINICAL STATUS.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# MACITENTAN

## Products Affected

- OPSUMIT

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	PULMONARY ARTERIAL HYPERTENSION (PAH): INITIAL: DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	PAH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# MARGETUXIMAB-CMKB

---

## Products Affected

- MARGENZA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# MARIBAVIR

---

## Products Affected

- LIVTENCITY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# MECASERMIN

## Products Affected

- INCRELEX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A ENDOCRINOLOGIST OR NEPHROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: OPEN EPIPHYSES AS CONFIRMED BY RADIOGRAPH OF WRIST AND HAND. RENEWAL: IMPROVEMENT WHILE ON THERAPY (I.E., INCREASE IN HEIGHT OR INCREASE IN HEIGHT VELOCITY).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# MECHLORETHAMINE

---

## Products Affected

- VALCHLOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# MEPOLIZUMAB

## Products Affected

- NUCALA SUBCUTANEOUS SOLUTION AUTO-INJECTOR
- NUCALA SUBCUTANEOUS SOLUTION RECONSTITUTED
- NUCALA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 100 MG/ML, 40 MG/0.4ML

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: ASTHMA: BLOOD EOSINOPHIL LEVEL OF AT LEAST 150 CELLS/MCL WITHIN THE PAST 12 MONTHS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: ASTHMA: PRESCRIBED BY OR IN CONSULTATION WITH A PHYSICIAN SPECIALIZING IN PULMONARY OR ALLERGY MEDICINE. CRSWNP: PRESCRIBED BY OR IN CONSULTATION WITH AN OTOLARYNGOLOGIST, ALLERGIST OR IMMUNOLOGIST.
<b>Coverage Duration</b>	INITIAL: ASTHMA: 12 MO. CRSWNP: 6 MO. OTHERS: 12 MO. RENEWAL: CRSWNP, ASTHMA: 12 MO.
<b>Other Criteria</b>	INITIAL: ASTHMA: 1) CONCURRENT THERAPY WITH A MEDIUM, HIGH-DOSE OR MAXIMALLY TOLERATED DOSE OF AN INHALED CORTICOSTEROID (ICS) AND AT LEAST ONE OTHER MAINTENANCE MEDICATION, 2) ONE OF THE FOLLOWING: (A) AT LEAST ONE ASTHMA EXACERBATION REQUIRING SYSTEMIC CORTICOSTEROID BURST LASTING 3 OR MORE DAYS WITHIN THE PAST 12 MONTHS OR AT LEAST ONE SERIOUS EXACERBATION REQUIRING HOSPITALIZATION OR ER VISIT WITHIN THE PAST 12 MONTHS, OR (B) POOR SYMPTOM CONTROL DESPITE CURRENT THERAPY AS EVIDENCED BY AT LEAST THREE OF THE FOLLOWING WITHIN THE PAST 4 WEEKS: DAYTIME ASTHMA SYMPTOMS MORE THAN TWICE/WEEK, ANY NIGHT WAKING DUE TO ASTHMA, SABA RELIEVER FOR SYMPTOMS MORE THAN TWICE/WEEK, ANY ACTIVITY LIMITATION DUE TO ASTHMA, AND 3) NO

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>CONCURRENT USE WITH XOLAIR, DUPIXENT, TEZSPIRE, OR OTHER ANTI-IL5 BIOLOGICS WHEN USED FOR ASTHMA. CRSWNP: 1) A 56 DAY TRIAL OF ONE TOPICAL NASAL CORTICOSTEROID, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR AN AUTOIMMUNE INDICATION. RENEWAL: ASTHMA: 1) NO CONCURRENT USE WITH XOLAIR, DUPIXENT, TEZSPIRE, OR OTHER ANTI-IL5 BIOLOGICS WHEN USED FOR ASTHMA, 2) CONTINUED USE OF ICS AND AT LEAST ONE OTHER MAINTENANCE MEDICATION, AND 3) CLINICAL RESPONSE AS EVIDENCED BY: (A) REDUCTION IN ASTHMA EXACERBATIONS FROM BASELINE, (B) DECREASED UTILIZATION OF RESCUE MEDICATIONS, (C) REDUCTION IN SEVERITY OR FREQUENCY OF ASTHMA-RELATED SYMPTOMS, OR (D) INCREASE IN PERCENT PREDICTED FEV1 FROM PRETREATMENT BASELINE. CRSWNP: 1) CLINICAL BENEFIT COMPARED TO BASELINE, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR AN AUTOIMMUNE INDICATION.</p>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# MIDOSTAURIN

## Products Affected

- RYDAPT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ACUTE MYELOID LEUKEMIA: 6 MONTHS. ADVANCED SYSTEMIC MASTOCYTOSIS: 12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# MIFEPRISTONE

## Products Affected

- *mifepristone oral tablet 300 mg*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	CUSHINGS SYNDROME (CS): INITIAL: DIAGNOSIS CONFIRMED BY: 1) 24-HR URINE FREE CORTISOL (2 OR MORE TESTS TO CONFIRM), 2) OVERNIGHT 1MG DEXAMETHASONE TEST, OR 3) LATE NIGHT SALIVARY CORTISOL (2 OR MORE TESTS TO CONFIRM).
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	CS: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS
<b>Other Criteria</b>	CS: INITIAL: HYPERCORTISOLISM IS NOT A RESULT OF CHRONIC GLUCOCORTICOIDS. RENEWAL: 1) CONTINUES TO HAVE IMPROVEMENT OF GLUCOSE TOLERANCE OR STABLE GLUCOSE TOLERANCE (E.G., REDUCED A1C, IMPROVED FASTING GLUCOSE, ETC.), 2) CONTINUES TO HAVE TOLERABILITY TO THERAPY, AND 3) CONTINUES TO NOT BE A CANDIDATE FOR SURGICAL TREATMENT OR HAS FAILED SURGERY.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# MILTEFOSINE

---

## Products Affected

- IMPAVIDO

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# MIRDAMETINIB

---

## Products Affected

- GOMEKLI ORAL CAPSULE 1 MG, 2 MG
- GOMEKLI ORAL TABLET SOLUBLE MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# MIRVETUXIMAB SORAVTANSINE-GYNX

## Products Affected

- ELAHERE

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER: AN OPHTHALMIC EXAM, INCLUDING VISUAL ACUITY AND SLIT LAMP EXAM, WILL BE COMPLETED PRIOR TO THE INITIATION OF THERAPY AND AT THE RECOMMENDED SCHEDULED INTERVALS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# MOMELOTINIB

---

## Products Affected

- OJAARA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# MOSUNETUZUMAB-AXGB

## Products Affected

- LUNSUMIO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA: INITIAL: 6 MONTHS. RENEWAL: 7 MONTHS.
Other Criteria	RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA: RENEWAL: 1) HAS ACHIEVED A PARTIAL RESPONSE TO TREATMENT, AND 2) HAS NOT PREVIOUSLY RECEIVED MORE THAN 17 CYCLES OF TREATMENT. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# NARCOLEPSY AGENTS

## Products Affected

- *armodafinil*
- *modafinil oral tablet 100 mg, 200 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All Medically-accepted Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# NAXITAMAB-GQGK

---

## Products Affected

- DANYELZA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# NERATINIB

## Products Affected

- NERLYNX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	EARLY-STAGE (STAGE I-III) BREAST CANCER: MEDICATION IS BEING REQUESTED WITHIN 2 YEARS OF COMPLETING THE LAST TRASTUZUMAB DOSE. ALL OTHER FDA APPROVED INDICATIONS ARE COVERED WITHOUT ADDITIONAL CRITERIA, EXCEPT THOSE CRITERIA IN THE FDA APPROVED LABEL.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# NILOTINIB

## Products Affected

- TASIGNA ORAL CAPSULE 150 MG, 200 MG, 50 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PREVIOUSLY TREATED PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (Ph+ CML): MUTATIONAL ANALYSIS PRIOR TO INITIATION AND TASIGNA IS APPROPRIATE PER NCCN GUIDELINE TABLE FOR TREATMENT RECOMMENDATIONS BASED ON BCR-ABL1 MUTATION PROFILE.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# NILOTINIB-DANZITEN

## Products Affected

- DANZITEN

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	PREVIOUSLY TREATED PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (Ph+ CML): 1) PERFORMED MUTATIONAL ANALYSIS PRIOR TO INITIATION OF THERAPY, AND 2) THERAPY IS APPROPRIATE PER NCCN GUIDELINE TABLE FOR TREATMENT RECOMMENDATIONS BASED ON BCR-ABL1 MUTATION PROFILE.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# NINTEDANIB

## Products Affected

- OFEV

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: IDIOPATHIC PULMONARY FIBROSIS (IPF): 1) A USUAL INTERSTITIAL PNEUMONIA (UIP) PATTERN AS EVIDENCED BY HIGH-RESOLUTION COMPUTED TOMOGRAPHY (HRCT) ALONE OR VIA A COMBINATION OF SURGICAL LUNG BIOPSY AND HRCT, AND 2) BASELINE FORCED VITAL CAPACITY (FVC) AT LEAST 50% OF PREDICTED VALUE. SYSTEMIC SCLEROSIS-ASSOCIATED INTERSTITIAL LUNG DISEASE (SSC-ILD): 1) AT LEAST 10% FIBROSIS ON A CHEST HRCT, AND 2) BASELINE FVC AT LEAST 40% OF PREDICTED VALUE. CHRONIC FIBROSING INTERSTITIAL LUNG DISEASE WITH A PROGRESSIVE PHENOTYPE (PF-ILD): 1) AT LEAST 10% FIBROSIS ON A CHEST HRCT, AND 2) BASELINE FVC AT LEAST 45% OF PREDICTED VALUE.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: IPF: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST. SSC-ILD, PF-ILD: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR RHEUMATOLOGIST.
<b>Coverage Duration</b>	INITIAL: SSC-ILD: 6 MOS. IPF, PF-ILD: 12 MOS. RENEWAL (ALL INDICATIONS): 12 MOS.
<b>Other Criteria</b>	INITIAL: IPF: 1) DOES NOT HAVE OTHER KNOWN CAUSES OF INTERSTITIAL LUNG DISEASE (E.G., CONNECTIVE TISSUE DISEASE, DRUG TOXICITY, ASBESTOS OR BERYLLIUM EXPOSURE, HYPERSENSITIVITY PNEUMONITIS), AND 2) TRIAL OF OR CONTRAINDICATION TO THE PREFERRED AGENT: ESBRIET (PIRFENIDONE). SSC-ILD: 1) DOES NOT HAVE OTHER KNOWN CAUSES OF INTERSTITIAL LUNG DISEASE (E.G., HEART FAILURE/FLUID OVERLOAD, DRUG-INDUCED LUNG TOXICITY, RECURRENT ASPIRATION), AND 2) TRIAL OF OR

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>CONTRAINDICATION TO THE PREFERRED AGENT: ACTEMRA SUBQ. PF-ILD: LUNG FUNCTION AND RESPIRATORY SYMPTOMS OR CHEST IMAGING HAVE WORSENERD/PROGRESSEED DESPITE TREATMENT WITH MEDICATIONS USED IN CLINICAL PRACTICE FOR ILD (NOT ATTRIBUTABLE TO COMORBIDITIES SUCH AS INFECTION, HEART FAILURE). RENEWAL: IPF, SSC-ILD, PF-ILD: CLINICAL MEANINGFUL IMPROVEMENT OR MAINTENANCE IN ANNUAL RATE OF DECLINE.</p>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# NIRAPARIB

## Products Affected

- ZEJULA ORAL CAPSULE
- ZEJULA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER: 1) ZEJULA WILL BE USED AS MONOTHERAPY, AND 2) ZEJULA IS STARTED NO LATER THAN 8 WEEKS AFTER THE MOST RECENT PLATINUM-CONTAINING REGIMEN.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# NIRAPARIB-ABIRATERONE

## Products Affected

- AKEEGA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC): 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# NIROGACESTAT

## Products Affected

- OGSIVEO ORAL TABLET 100 MG, 150 MG, 50 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# NITISINONE

## Products Affected

- *nitisinone*
- ORFADIN ORAL SUSPENSION

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	HEREDITARY TYROSINEMIA TYPE 1 (HT-1): INITIAL: DIAGNOSIS CONFIRMED BY ELEVATED URINARY OR PLASMA SUCCINYLACETONE LEVELS OR A MUTATION IN THE FUMARYLACETOACETATE HYDROLASE GENE. RENEWAL: URINARY OR PLASMA SUCCINYLACETONE LEVELS HAVE DECREASED FROM BASELINE WHILE ON TREATMENT WITH NITISINONE.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	HT-1: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A PRESCRIBER SPECIALIZING IN INHERITED METABOLIC DISEASES.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	HT-1: INITIAL: ORFADIN SUSPENSION: TRIAL OF OR CONTRAINDICATION TO PREFERRED NITISINONE TABLETS OR CAPSULES.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# NIVOLUMAB

---

## Products Affected

- OPDIVO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	UNRESECTABLE OR METASTATIC MELANOMA: NO CONCURRENT USE WITH TARGETED THERAPY (I.E., BRAF INHIBITORS, MEK INHIBITORS, AND NTRK INHIBITORS).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# NIVOLUMAB-HYALURONIDASE-NVHY

## Products Affected

- OPDIVO QVANTIG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# NIVOLUMAB-RELATLIMAB-RMBW

---

## Products Affected

- OPDUALAG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# NOGAPENDEKIN ALFA

---

## Products Affected

- ANKTIVA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	40 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# OCRELIZUMAB

## Products Affected

- OCREVUS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	RELAPSING FORM OF MULTIPLE SCLEROSIS (MS): TRIAL OF TWO AGENTS INDICATED FOR THE TREATMENT OF RELAPSING FORMS OF MS. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# OCRELIZUMAB-HYALURONIDASE-OCSQ

## Products Affected

- OCREVUS ZUNOVO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	RELAPSING FORM OF MULTIPLE SCLEROSIS (MS): TRIAL OF TWO AGENTS INDICATED FOR THE TREATMENT OF RELAPSING FORMS OF MS. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# OFATUMUMAB-SQ

## Products Affected

- KESIMPTA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# OLANZAPINE/SAMIDORPHAN

## Products Affected

- LYBALVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	SCHIZOPHRENIA, BIPOLAR I: PRESCRIBED BY OR IN CONSULTATION WITH A PSYCHIATRIST
Coverage Duration	12 MONTHS
Other Criteria	SCHIZOPHRENIA: 1) AT HIGH RISK FOR WEIGHT GAIN, AND 2) TRIAL OF OR CONTRAINDICATION TO A FORMULARY VERSION OF LURASIDONE OR ONE OF THE FOLLOWING ORAL ANTIPSYCHOTICS: RISPERIDONE, CLOZAPINE TABLET, OLANZAPINE, IMMEDIATE RELEASE QUETIAPINE FUMARATE, ZIPRASIDONE, ARIPIPRAZOLE. BIPOLAR I: 1) AT HIGH RISK FOR WEIGHT GAIN, AND 2) TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING ORAL ANTIPSYCHOTICS: RISPERIDONE, OLANZAPINE, IMMEDIATE RELEASE QUETIAPINE FUMARATE, ZIPRASIDONE, ARIPIPRAZOLE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# OLAPARIB

## Products Affected

- LYNPARZA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER: MEDICATION WILL BE USED AS MONOTHERAPY. METASTATIC CASTRATION-RESISTANT PROSTATE CANCER: 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG. ALL OTHER FDA APPROVED INDICATIONS ARE COVERED WITHOUT ADDITIONAL CRITERIA, EXCEPT THOSE CRITERIA IN THE FDA APPROVED LABEL.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# OLUTASIDENIB

---

## Products Affected

- REZLIDHIA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# OMACETAXINE

---

## Products Affected

- SYNRIPO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# OMALIZUMAB

## Products Affected

- XOLAIR

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: ASTHMA: POSITIVE SKIN PRICK OR BLOOD TEST (E.G., ELISA, FEIA) TO A PERENNIAL AEROALLERGEN AND A BASELINE IGE SERUM LEVEL OF AT LEAST 30 IU/ML. FOOD ALLERGY: 1) IGE SERUM LEVEL OF AT LEAST 30 IU/ML, AND 2) ALLERGEN SPECIFIC IGE SERUM LEVEL OF AT LEAST 6 KUA/L TO AT LEAST ONE FOOD, OR POSITIVE SKIN PRICK TEST TO AT LEAST ONE FOOD, OR POSITIVE MEDICALLY MONITORED FOOD CHALLENGE TO AT LEAST ONE FOOD.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL/RENEWAL: CHRONIC SPONTANEOUS URTICARIA (CSU): PRESCRIBED BY OR IN CONSULTATION WITH AN ALLERGIST, DERMATOLOGIST, OR IMMUNOLOGIST. INITIAL: CHRONIC RHINOSINUSITIS WITH NASAL POLYPS (CRSWNP): PRESCRIBED BY OR IN CONSULTATION WITH AN OTOLARYNGOLOGIST, ALLERGIST OR IMMUNOLOGIST. ASTHMA: PRESCRIBED BY OR IN CONSULTATION WITH A PHYSICIAN SPECIALIZING IN ALLERGY OR PULMONARY MEDICINE. FOOD ALLERGY: PRESCRIBED BY OR IN CONSULTATION WITH ALLERGIST OR IMMUNOLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: ASTHMA 12 MO/12 MO, CSU 6 MO/12 MO, CRSWNP 6 MO/12 MO, FOOD ALLERGY 12 MO/24 MO
<b>Other Criteria</b>	INITIAL: CSU: 1) TRIAL OF AND MAINTAINED ON, OR CONTRAINDICATION TO A SECOND GENERATION H1 ANTI-HISTAMINE AND 2) STILL EXPERIENCES HIVES OR ANGIOEDEMA ON MOST DAYS OF THE WEEK FOR AT LEAST 6 WEEKS. CRSWNP: 1) A 56 DAY TRIAL OF ONE TOPICAL NASAL CORTICOSTEROID, 2) TRIAL OF OR CONTRAINDICATION TO ONE PREFERRED AGENT: NUCALA, DUPIXENT, AND 3) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR AN AUTOIMMUNE INDICATION. ASTHMA: 1) CONCURRENT THERAPY WITH A MEDIUM, HIGH-DOSE OR MAXIMALLY TOLERATED DOSE OF AN INHALED CORTICOSTEROID (ICS) AND AT LEAST ONE OTHER MAINTENANCE MEDICATION, 2) ONE OF THE FOLLOWING: (A) AT LEAST ONE ASTHMA EXACERBATION REQUIRING SYSTEMIC CORTICOSTEROID BURST LASTING 3 OR MORE DAYS WITHIN THE PAST 12 MONTHS OR AT LEAST ONE SERIOUS EXACERBATION REQUIRING HOSPITALIZATION OR ER VISIT WITHIN THE PAST 12 MONTHS, OR (B) POOR SYMPTOM CONTROL DESPITE CURRENT THERAPY AS EVIDENCED BY AT LEAST THREE OF THE FOLLOWING WITHIN THE PAST 4 WEEKS: DAYTIME ASTHMA SYMPTOMS MORE THAN TWICE/WEEK, ANY NIGHT WAKING DUE TO ASTHMA, SABA RELIEVER FOR SYMPTOMS MORE THAN TWICE/WEEK, ANY ACTIVITY LIMITATION DUE TO ASTHMA, AND 3) NO CONCURRENT USE WITH DUPIXENT, TEZSPIRE, OR ANTI-IL5 BIOLOGICS WHEN USED FOR ASTHMA. FOOD ALLERGY: 1) CONCURRENT USE WITH AN ACTIVE PRESCRIPTION FOR EPINEPHRINE AUTO-INJECTOR/INJECTION, AND 2) NO CONCURRENT USE WITH PEANUT-SPECIFIC IMMUNOTHERAPY. RENEWAL: CSU: MAINTAINED ON OR CONTRAINDICATION TO A SECOND GENERATION H1 ANTI-HISTAMINE. CRSWNP: 1) CLINICAL BENEFIT COMPARED TO BASELINE, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR AN AUTOIMMUNE INDICATION. ASTHMA: 1) NO CONCURRENT USE WITH DUPIXENT, TEZSPIRE, OR ANTI-IL5 BIOLOGICS WHEN USED FOR ASTHMA, 2) CONTINUED USE OF ICS AND AT LEAST ONE OTHER MAINTENANCE MEDICATION, AND 3) CLINICAL RESPONSE AS EVIDENCED BY ONE OF THE FOLLOWING: (A) REDUCTION IN ASTHMA EXACERBATIONS FROM BASELINE, (B) DECREASED UTILIZATION OF RESCUE MEDICATIONS, (C) REDUCTION IN SEVERITY OR FREQUENCY OF ASTHMA-RELATED SYMPTOMS, OR (D) INCREASE IN PERCENT PREDICTED FEV1 FROM PRETREATMENT BASELINE. FOOD ALLERGY: 1) PERSISTENT IGE-MEDIATED FOOD ALLERGY, 2) CONCURRENT USE WITH AN ACTIVE PRESCRIPTION FOR EPINEPHRINE AUTO-INJECTOR/INJECTION, AND 3) NO</p>

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
	CONCURRENT USE WITH PEANUT-SPECIFIC IMMUNOTHERAPY.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# OSIMERTINIB

---

## Products Affected

- TAGRISSO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# OXANDROLONE

## Products Affected

- *oxandrolone oral*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 MONTHS
Other Criteria	PROTEIN CATABOLISM, BONE PAIN: 1) MONITORED FOR PELIOSIS HEPATIS, LIVER CELL TUMORS, AND BLOOD LIPID CHANGES, 2) DOES NOT HAVE KNOWN OR SUSPECTED: CARCINOMA OF THE PROSTATE OR BREAST IN MALE PATIENTS, CARCINOMA OF THE BREAST IN FEMALES WITH HYPERCALCEMIA, NEPHROSIS (THE NEPHROTIC PHASE OF NEPHRITIS), OR HYPERCALCEMIA, AND 3) DOES NOT HAVE SEVERE HEPATIC DYSFUNCTION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# PACRITINIB

---

## Products Affected

- VONJO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	MYELOFIBROSIS: RENEWAL: CONTINUES TO BENEFIT FROM THE MEDICATION
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# PALBOCICLIB

## Products Affected

- IBRANCE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	ADVANCED OR METASTATIC BREAST CANCER: TRIAL OF OR CONTRAINDICATION TO ONE OF THE PREFERRED AGENTS, WHERE INDICATIONS ALIGN: KISQALI, VERZENIO.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# PARATHYROID HORMONE

## Products Affected

- NATPARA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	HYPOCALCEMIA SECONDARY TO HYPOPARATHYROIDISM: PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	HYPOCALCEMIA SECONDARY TO HYPOPARATHYROIDISM: 1) TRIAL OF OR CONTRAINDICATION TO CALCITRIOL, 2) HYPOPARATHYROIDISM IS NOT DUE TO A CALCIUM SENSING RECEPTOR (CSR) MUTATION, AND 3) HYPOPARATHYROIDISM IS NOT CONSIDERED ACUTE POST-SURGICAL HYPOPARATHYROIDISM.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# PASIREOTIDE DIASPARTATE

## Products Affected

- SIGNIFOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	CUSHINGS DISEASE (CD): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS.
Other Criteria	CD: RENEWAL: 1) CONTINUED IMPROVEMENT OF CUSHINGS DISEASE, AND 2) MAINTAINED TOLERABILITY TO SIGNIFOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# PAZOPANIB

## Products Affected

- *pazopanib hcl*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	ADVANCED SOFT TISSUE SARCOMA (STS): NOT USED FOR ADIPOCYTIC STS OR GASTROINTESTINAL STROMAL TUMORS (GIST)
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# PEGFILGRASTIM - APGF

## Products Affected

- NYVEPRIA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# PEGFILGRASTIM-NEULASTA ONPRO

---

## Products Affected

- NEULASTA ONPRO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# PEGINTERFERON ALFA-2A

## Products Affected

- PEGASYS SUBCUTANEOUS SOLUTION 180 MCG/ML
- PEGASYS SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	HEPATITIS B: PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST, INFECTIOUS DISEASE SPECIALIST, OR PHYSICIAN SPECIALIZING IN THE TREATMENT OF HEPATITIS (E.G., HEPATOLOGIST).
Coverage Duration	HEP B/HEP C: 48 WEEKS.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# PEGVISOMANT

## Products Affected

- SOMAVERT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# PEMBROLIZUMAB

## Products Affected

- KEYTRUDA INTRAVENOUS SOLUTION

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	UNRESECTABLE OR METASTATIC MELANOMA: NO CONCURRENT USE WITH TARGETED THERAPY (I.E., BRAF INHIBITORS, MEK INHIBITORS, AND NTRK INHIBITORS).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# PEMIGATINIB

## Products Affected

- PEMAZYRE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	CHOLANGIOCARCINOMA, MYELOID/LYMPHOID NEOPLASMS: COMPREHENSIVE OPHTHALMOLOGICAL EXAMINATION, INCLUDING OPTICAL COHERENCE TOMOGRAPHY (OCT), WILL BE COMPLETED PRIOR TO INITIATION OF THERAPY AND AT THE RECOMMENDED SCHEDULED INTERVALS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# PENICILLAMINE TABLET

## Products Affected

- *penicillamine oral tablet*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: CYSTINURIA: HAS NEPHROLITHIASIS AND ONE OF THE FOLLOWING: 1) STONE ANALYSIS SHOWING PRESENCE OF CYSTINE, 2) PRESENCE OF PATHOGNOMONIC HEXAGONAL CYSTINE CRYSTALS ON URINALYSIS, OR 3) FAMILY HISTORY OF CYSTINURIA AND POSITIVE CYANIDE-NITROPRUSSIDE SCREENING.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: WILSONS DISEASE: PRESCRIBED BY OR IN CONSULTATION WITH A HEPATOLOGIST OR GASTROENTEROLOGIST. CYSTINURIA: PRESCRIBED BY OR IN CONSULTATION WITH A NEPHROLOGIST. RHEUMATOID ARTHRITIS (RA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST.
<b>Coverage Duration</b>	INITIAL: 12 MONTHS, RENEWAL: LIFETIME.
<b>Other Criteria</b>	INITIAL: WILSONS DISEASE: 1) LEIPZIG SCORE OF 4 OR GREATER. RA: 1) NO HISTORY OR OTHER EVIDENCE OF RENAL INSUFFICIENCY, AND 2) TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. RENEWAL: RA: 1) NO HISTORY OR OTHER EVIDENCE OF RENAL INSUFFICIENCY, AND 2) EXPERIENCED OR MAINTAINED IMPROVEMENT IN TENDER JOINT COUNT OR SWOLLEN JOINT COUNT COMPARED TO BASELINE. WILSONS DISEASE, CYSTINURIA: CONTINUES TO BENEFIT FROM THE MEDICATION.
<b>Indications</b>	All FDA-approved Indications.

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# PEXIDARTINIB

---

## Products Affected

- TURALIO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# PIMAVANSERIN

## Products Affected

- NUPLAZID ORAL CAPSULE
- NUPLAZID ORAL TABLET 10 MG

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	PSYCHOSIS IN PARKINSONS DISEASE (PD): INITIAL: 18 YEARS OR OLDER
<b>Prescriber Restrictions</b>	PSYCHOSIS IN PD: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST, GERIATRICIAN, OR A BEHAVIORAL HEALTH SPECIALIST (E.G., PSYCHIATRIST).
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	PSYCHOSIS IN PD: RENEWAL: IMPROVEMENT IN PSYCHOSIS SYMPTOMS FROM BASELINE AND DEMONSTRATES A CONTINUED NEED FOR TREATMENT.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# PIRFENIDONE

## Products Affected

- *pirfenidone oral capsule*
- *pirfenidone oral tablet 267 mg, 534 mg, 801 mg*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	IDIOPATHIC PULMONARY FIBROSIS (IPF): INITIAL: 1) A USUAL INTERSTITIAL PNEUMONIA (UIP) PATTERN AS EVIDENCED BY HIGH-RESOLUTION COMPUTED TOMOGRAPHY (HRCT) ALONE OR VIA A COMBINATION OF SURGICAL LUNG BIOPSY AND HRCT, AND 2) PREDICTED FORCED VITAL CAPACITY (FVC) OF AT LEAST 50% AT BASELINE.
<b>Age Restrictions</b>	IPF: INITIAL: 18 YEARS OR OLDER.
<b>Prescriber Restrictions</b>	IPF: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	IPF: INITIAL: 1) DOES NOT HAVE OTHER KNOWN CAUSES OF INTERSTITIAL LUNG DISEASE (E.G., CONNECTIVE TISSUE DISEASE, DRUG TOXICITY, ASBESTOS OR BERYLLIUM EXPOSURE, HYPERSENSITIVITY PNEUMONITIS, SYSTEMIC SCLEROSIS, RHEUMATOID ARTHRITIS, RADIATION, SARCOIDOSIS, BRONCHIOLITIS OBLITERANS ORGANIZING PNEUMONIA, HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION, VIRAL HEPATITIS, OR CANCER). RENEWAL: CLINICAL MEANINGFUL IMPROVEMENT OR MAINTENANCE IN ANNUAL RATE OF DECLINE.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# PIRTOBRUTINIB

---

## Products Affected

- JAYPIRCA ORAL TABLET 100 MG, 50 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# POMALIDOMIDE

---

## Products Affected

- POMALYST

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# PONATINIB

---

## Products Affected

- ICLUSIG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	CHRONIC MYELOID LEUKEMIA (CML): MUTATIONAL ANALYSIS PRIOR TO INITIATION AND ICLUSIG IS APPROPRIATE PER NCCN GUIDELINE TABLE FOR TREATMENT RECOMMENDATIONS BASED ON BCR-ABL1 MUTATION PROFILE.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# POSACONAZOLE TABLET

## Products Affected

- *posaconazole oral tablet delayed release*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	CONTINUATION OF THERAPY AFTER HOSPITAL DISCHARGE, PROPHYLAXIS: 6 MONTHS. TREATMENT: 12 WEEKS.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# PRALSETINIB

---

## Products Affected

- GAVRETO

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# PYRIMETHAMINE

## Products Affected

- *pyrimethamine oral*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	TOXOPLASMOSIS: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH AN INFECTIOUS DISEASE SPECIALIST.
Coverage Duration	TOXOPLASMOSIS: INITIAL: 8 WEEKS, RENEWAL: 6 MOS.
Other Criteria	TOXOPLASMOSIS: RENEWAL: ONE OF THE FOLLOWING: (1) PERSISTENT CLINICAL DISEASE (HEADACHE, NEUROLOGICAL SYMPTOMS, OR FEVER) AND PERSISTENT RADIOGRAPHIC DISEASE (ONE OR MORE MASS LESIONS ON BRAIN IMAGING), OR (2) CD4 COUNT LESS THAN 200 CELLS/MM3 AND CURRENTLY TAKING AN ANTI-RETROVIRAL THERAPY IF HIV POSITIVE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# QUININE

---

## Products Affected

- *quinine sulfate oral*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# QUIZARTINIB

---

## Products Affected

- VANFLYTA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# REGORAFENIB

---

## Products Affected

- STIVARGA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# RELUGOLIX

---

## Products Affected

- ORGOVYX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# REPOTRECTINIB

---

## Products Affected

- AUGTYRO ORAL CAPSULE 160 MG, 40 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# RESLIZUMAB

## Products Affected

- CINQAIR

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	ASTHMA: INITIAL: BLOOD EOSINOPHIL LEVEL OF AT LEAST 150 CELLS/MCL WITHIN THE PAST 12 MONTHS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	ASTHMA: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A PHYSICIAN SPECIALIZING IN ALLERGY OR PULMONARY MEDICINE.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	ASTHMA: INITIAL: 1) CONCURRENT THERAPY WITH A MEDIUM, HIGH-DOSE OR MAXIMALLY TOLERATED DOSE OF AN INHALED CORTICOSTEROID (ICS) AND ONE OTHER MAINTENANCE MEDICATION, 2) ONE ASTHMA EXACERBATION REQUIRING SYSTEMIC CORTICOSTEROID BURST LASTING 3 OR MORE DAYS WITHIN THE PAST 12 MONTHS, OR AT LEAST ONE SERIOUS EXACERBATION REQUIRING HOSPITALIZATION OR ER VISIT WITHIN THE PAST 12 MONTHS, OR POOR SYMPTOM CONTROL DESPITE CURRENT THERAPY AS EVIDENCED BY AT LEAST THREE OF THE FOLLOWING WITHIN THE PAST 4 WEEKS: (A) DAYTIME ASTHMA SYMPTOMS MORE THAN TWICE/WEEK, (B) ANY NIGHT WAKING DUE TO ASTHMA, (C) SABA RELIEVER FOR SYMPTOMS MORE THAN TWICE/WEEK, (D) ANY ACTIVITY LIMITATION DUE TO ASTHMA, 3) TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: FASENRA, NUCALA, DUPIXENT, AND 4) NO CONCURRENT USE WITH XOLAIR, DUPIXENT, TEZSPIRE, OR OTHER ANTI-IL5 BIOLOGICS WHEN USED FOR ASTHMA. RENEWAL: 1) NO CONCURRENT USE WITH XOLAIR, DUPIXENT,

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	TEZSPIRE, OR OTHER ANTI-IL5 BIOLOGICS WHEN USED FOR ASTHMA, 2) CONTINUED USE OF ICS AND ONE OTHER MAINTENANCE MEDICATION, AND 3) CLINICAL RESPONSE AS EVIDENCED BY: (A) REDUCTION IN ASTHMA EXACERBATIONS FROM BASELINE, (B) DECREASED UTILIZATION OF RESCUE MEDICATIONS, (C) REDUCTION IN SEVERITY OR FREQUENCY OF ASTHMA-RELATED SYMPTOMS, OR (D) INCREASE IN PERCENT PREDICTED FEV1 FROM PRETREATMENT BASELINE. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# RETIFANLIMAB-DLWR

## Products Affected

- ZYNYZ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# REVUMENIB

---

## Products Affected

- REVUFORJ ORAL TABLET 110 MG, 160 MG, 25 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# RIBOCICLIB

## Products Affected

- KISQALI (200 MG DOSE)
- KISQALI (400 MG DOSE)
- KISQALI (600 MG DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# RIBOCICLIB-LETROZOLE

## Products Affected

- KISQALI FEMARA (200 MG DOSE)
- KISQALI FEMARA (400 MG DOSE)
- KISQALI FEMARA (600 MG DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# RIFAXIMIN

## Products Affected

- XIFAXAN ORAL TABLET 200 MG, 550 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	TRAVELERS DIARRHEA, HEPATIC ENCEPHALOPATHY (HE): 12 MOS. IBS-D: 8 WKS.
Other Criteria	HE: TRIAL OF OR CONTRAINDICATION TO LACTULOSE OR CONCURRENT LACTULOSE THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# RILONACEPT

## Products Affected

- ARCALYST

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	<p>CRYOPYRIN-ASSOCIATED PERIODIC SYNDROMES (CAPS): 1) ONE OF THE FOLLOWING: (A) GENETIC TEST FOR GAIN-OF-FUNCTION MUTATIONS IN THE NLRP3 GENE, OR (B) HAS INFLAMMATORY MARKERS (I.E., ELEVATED CRP, ESR, SERUM AMYLOID A PROTEIN (SAA) OR S100 PROTEINS), AND 2) TWO OF THE FOLLOWING: URTICARIAL-LIKE RASH (NEUTROPHILIC DERMATITIS), COLD-TRIGGERED EPISODES, SENSORINEURAL HEARING LOSS, MUSCULOSKELETAL SYMPTOMS, CHRONIC ASEPTIC MENINGITIS, SKELETAL ABNORMALITIES.</p> <p>DEFICIENCY OF INTERLEUKIN-1 RECEPTOR ANTAGONIST (DIRA): 1) ONE OF THE FOLLOWING: (A) GENETIC TEST FOR GAIN-OF-FUNCTION MUTATIONS IN THE IL1RN GENE, OR (B) HAS INFLAMMATORY MARKERS (I.E., ELEVATED CRP, ESR), AND 2) ONE OF THE FOLLOWING: PUSTULAR PSORIASIS-LIKE RASHES, OSTEOMYELITIS, ABSENCE OF BACTERIAL OSTEOMYELITIS, ONYCHOMADESIS. RECURRENT PERICARDITIS (RP): TWO OF THE FOLLOWING: CHEST PAIN CONSISTENT WITH PERICARDITIS, PERICARDIAL FRICTION RUB, ECG SHOWING DIFFUSE ST-SEGMENT ELEVATION OR PR-SEGMENT DEPRESSION, NEW OR WORSENING PERICARDIAL EFFUSION.</p>
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	CAPS, DIRA: LIFETIME. RP: 12 MONTHS.
<b>Other Criteria</b>	<p>CAPS: NO CONCURRENT USE WITH OTHER IL-1 INHIBITORS.</p> <p>DIRA: 1) NO CONCURRENT USE WITH OTHER IL-1 INHIBITORS, AND 2) TRIAL OF THE PREFERRED AGENT: KINERET. RP: 1)</p>

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
	HAD AN EPISODE OF ACUTE PERICARDITIS, 2) SYMPTOM-FREE FOR 4 TO 6 WEEKS, AND 3) NO CONCURRENT USE WITH OTHER IL-1 INHIBITORS.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# RILUZOLE

---

## Products Affected

- TIGLUTIK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OR OLDER
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	AMYOTROPHIC LATERAL SCLEROSIS (ALS): (1) TRIAL OF RILUZOLE TABLETS, AND (2) PATIENT IS UNABLE TO TAKE TABLET FORMULATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# RIMEGEPANT

## Products Affected

- NURTEC

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	<p>INITIAL: ACUTE MIGRAINE TREATMENT: 1) TRIAL OF OR CONTRAINDICATION TO ONE TRIPTAN (E.G., SUMATRIPTAN, RIZATRIPTAN), AND 2) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR ACUTE MIGRAINE TREATMENT.</p> <p>EPISODIC MIGRAINE PREVENTION: 1) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR MIGRAINE PREVENTION, AND 2) TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING PREVENTIVE MIGRAINE TREATMENTS: DIVALPROEX SODIUM, TOPIRAMATE, PROPRANOLOL, TIMOLOL. RENEWAL: ACUTE MIGRAINE TREATMENT: 1) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR ACUTE MIGRAINE TREATMENT, AND 2) ONE OF THE FOLLOWING: (A) IMPROVEMENT FROM BASELINE IN A VALIDATED ACUTE TREATMENT PATIENT-REPORTED OUTCOME QUESTIONNAIRE, OR (B) THERAPY WORKS CONSISTENTLY IN MAJORITY OF MIGRAINE ATTACKS.</p> <p>EPISODIC MIGRAINE PREVENTION: 1) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR MIGRAINE PREVENTION, AND 2) REDUCTION IN MIGRAINE OR HEADACHE FREQUENCY, MIGRAINE SEVERITY, OR MIGRAINE DURATION WITH THERAPY.</p>

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# RIOCIGUAT

## Products Affected

- ADEMPAS

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PULMONARY ARTERIAL HYPERTENSION (PAH): DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS. PERSISTENT/RECURRENT CHRONIC THROMBOEMBOLIC PULMONARY HYPERTENSION (CTEPH) (WHO GROUP 4): WHO FUNCTIONAL CLASS II-IV SYMPTOMS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: PAH, CTEPH: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: PAH: NO CONCURRENT USE WITH NITRATES, NITRIC OXIDE DONORS, PHOSPHODIESTERASE (PDE) INHIBITORS, OR NON-SPECIFIC PDE INHIBITORS. CTEPH: 1) NO CONCURRENT USE WITH NITRATES, NITRIC OXIDE DONORS, PDE INHIBITORS, OR NON-SPECIFIC PDE INHIBITORS, AND 2) NOT A CANDIDATE FOR SURGERY OR HAS INOPERABLE CTEPH OR HAS PERSISTENT OR RECURRENT DISEASE AFTER SURGICAL TREATMENT. RENEWAL: PAH, CTEPH: NO CONCURRENT USE WITH NITRATES, NITRIC OXIDE DONORS, PDE INHIBITORS, OR NON-SPECIFIC PDE INHIBITORS.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# RIPRETINIB

---

## Products Affected

- QINLOCK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# RISANKIZUMAB-RZAA

## Products Affected

- SKYRIZI
- SKYRIZI (150 MG DOSE)
- SKYRIZI PEN

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PLAQUE PSORIASIS (PSO): PLAQUE PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP, OR GENITAL AREA.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
<b>Other Criteria</b>	INITIAL: PSO: 1) ONE OF THE FOLLOWING: (A) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY) FOR THE TREATMENT OF PSO, (B) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA FOR THE TREATMENT OF PSO, OR (C) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PSO. PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>TARGETED SMALL MOLECULES FOR UC. RENEWAL: PSO: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC.</p>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# RISDIPLAM

## Products Affected

- EVRYSDI ORAL SOLUTION RECONSTITUTED

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	SPINAL MUSCULAR ATROPHY (SMA): INITIAL: GENE MUTATION ANALYSIS INDICATING MUTATIONS OR DELETIONS OF BOTH ALLELES OF THE SURVIVAL MOTOR NEURON 1 (SMN1) GENE. FOR PRESYMPTOMATIC PATIENTS: UP TO THREE COPIES OF SURVIVAL MOTOR NEURON 2 (SMN2) BASED ON NEWBORN SCREENING.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	SMA: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROMUSCULAR SPECIALIST OR SMA SPECIALIST AT A SMA SPECIALTY CENTER.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS
<b>Other Criteria</b>	SMA: INITIAL: FOR SYMPTOMATIC PATIENTS: 1) BASELINE MOTOR FUNCTION ASSESSMENT BY A NEUROMUSCULAR SPECIALIST OR SMA SPECIALIST, AND 2) IF PATIENT RECEIVED GENE THERAPY, PATIENT HAD LESS THAN EXPECTED CLINICAL BENEFIT WITH GENE THERAPY. RENEWAL: IMPROVED, MAINTAINED, OR DEMONSTRATED LESS THAN EXPECTED DECLINE IN: 1) MOTOR FUNCTION ASSESSMENTS COMPARED TO BASELINE, OR 2) OTHER MUSCLE FUNCTION.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# RITUXIMAB AND HYALURONIDASE HUMAN-SQ

## Products Affected

- RITUXAN HYCELA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	FOLLICULAR LYMPHOMA (FL), DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL), CHRONIC LYMPHOCYTIC LEUKEMIA (CLL): HAS RECEIVED OR WILL RECEIVE AT LEAST ONE FULL DOSE OF A RITUXIMAB PRODUCT BY INTRAVENOUS INFUSION PRIOR TO INITIATION OF RITUXIMAB AND HYALURONIDASE. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# RITUXIMAB-ABBS

## Products Affected

- TRUXIMA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. NON-HODGKINS LYMPHOMA (NHL), CHRONIC LYMPHOCYTIC LEUKEMIA (CLL): PRESCRIBED BY OR IN CONSULTATION WITH AN ONCOLOGIST.
Coverage Duration	RA: INITIAL: 6 MO, RENEWAL: 12 MO. NHL, GPA, MPA: 12 MO. CLL: 6 MO.
Other Criteria	RA: INITIAL: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ, RINVOQ, ORENCIA. RENEWAL: CONTINUES TO BENEFIT FROM THE MEDICATION. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# RITUXIMAB-ARRX

## Products Affected

- RIABNI

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	RHEUMATOID ARTHRITIS (RA): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. NON-HODGKINS LYMPHOMA (NHL), CHRONIC LYMPHOCYTIC LEUKEMIA (CLL): PRESCRIBED BY OR IN CONSULTATION WITH AN ONCOLOGIST.
<b>Coverage Duration</b>	RA: INITIAL: 6 MO, RENEWAL: 12 MO. NHL, GPA, MPA: 12 MO. CLL: 6 MO.
<b>Other Criteria</b>	RA: INITIAL: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ, RINVOQ, ORENCIA. RENEWAL: CONTINUES TO BENEFIT FROM THE MEDICATION. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# RITUXIMAB-PVVR

## Products Affected

- RUXIENCE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	RA: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. NON-HODGKINS LYMPHOMA (NHL), CHRONIC LYMPHOCYTIC LEUKEMIA (CLL): PRESCRIBED BY OR IN CONSULTATION WITH AN ONCOLOGIST.
Coverage Duration	RA: INITIAL: 6 MO, RENEWAL: 12 MO. NHL, GPA, MPA: 12 MO. CLL: 6 MO.
Other Criteria	RA: INITIAL: TRIAL OF OR CONTRAINDICATION TO TWO PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ, RINVOQ, ORENCIA. RENEWAL: CONTINUES TO BENEFIT FROM THE MEDICATION. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# ROPEGINTERFERON ALFA-2B-NJFT

---

## Products Affected

- BESREMI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# RUCAPARIB

## Products Affected

- RUBRACA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC CASTRATION-RESISTANT PROSTATE CANCER: ONE OF THE FOLLOWING: 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# RUXOLITINIB

## Products Affected

- JAKAFI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	MYELOFIBROSIS: INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS. POLYCYTHEMIA VERA, GVHD: 12 MONTHS.
Other Criteria	MYELOFIBROSIS: RENEWAL: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# SAPROPTERIN

## Products Affected

- *javygtor oral tablet*
- *sapropterin dihydrochloride oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 2 MONTHS, RENEWAL 12 MONTHS.
Other Criteria	HYPERPHENYLALANINEMIA (HPA): INITIAL: NO CONCURRENT USE WITH PALYNZIQ. RENEWAL: 1) CONTINUES TO BENEFIT FROM TREATMENT, AND 2) NO CONCURRENT USE WITH PALYNZIQ.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# SECUKINUMAB IV

## Products Affected

- COSENTYX INTRAVENOUS

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: NON-RADIOGRAPHIC AXIAL SPONDYLOARTHRITIS (NR-AXSPA): 1) C-REACTIVE PROTEIN (CRP) LEVELS ABOVE THE UPPER LIMIT OF NORMAL, OR 2) SACROILIITIS ON MAGNETIC RESONANCE IMAGING (MRI).
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR A DERMATOLOGIST. ANKYLOSING SPONDYLITIS (AS), NR-AXSPA: PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
<b>Other Criteria</b>	INITIAL: PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PSA. AS: 1) TRIAL OF OR CONTRAINDICATION TO AN NSAID, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. NR-AXSPA: 1) TRIAL OF OR CONTRAINDICATION TO AN NSAID, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR NR-AXSPA. RENEWAL: PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. AS: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. NR-AXSPA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
	TARGETED SMALL MOLECULES FOR NR-AXSPA. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# SECUKINUMAB SQ

## Products Affected

- COSENTYX (300 MG DOSE)
- COSENTYX SENSOREADY (300 MG)
- COSENTYX SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 75 MG/0.5ML
- COSENTYX UNOREADY

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP, OR FACE. NON-RADIOGRAPHIC AXIAL SPONDYLOARTHRITIS (NR-AXSPA): 1) C-REACTIVE PROTEIN (CRP) LEVELS ABOVE THE UPPER LIMIT OF NORMAL, OR 2) SACROILIITIS ON MAGNETIC RESONANCE IMAGING (MRI).
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: PSO, HIDRADENITIS SUPPURATIVA (HS): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR A DERMATOLOGIST. ANKYLOSING SPONDYLITIS (AS), NR-AXSPA, ENTHESITIS-RELATED ARTHRITIS (ERA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST.
<b>Coverage Duration</b>	INITIAL: HS: 12 MONTHS, ALL OTHER INDICATIONS: 6 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: PSO: 1) ONE OF THE FOLLOWING: (A) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY) FOR THE TREATMENT OF PSO, (B) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA FOR THE TREATMENT OF PSO, OR (C) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION, AND 2) NO CONCURRENT USE WITH

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PSO. PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. AS: 1) TRIAL OF OR CONTRAINDICATION TO AN NSAID, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. NR-AXSPA: 1) TRIAL OF OR CONTRAINDICATION TO AN NSAID, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR NR-AXSPA. ERA: TRIAL OF OR CONTRAINDICATION TO ONE NSAID, SULFASALAZINE, OR METHOTREXATE. HS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR HS. RENEWAL: PSO: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. AS: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. NR-AXSPA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR NR-AXSPA. ERA, HS: CONTINUES TO BENEFIT FROM THE MEDICATION.</p>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# SELEXIPAG

## Products Affected

- UPTRAVI INTRAVENOUS
- UPTRAVI ORAL TABLET 1000 MCG, 1200 MCG, 1400 MCG, 1600 MCG, 200 MCG, 400 MCG, 600 MCG, 800 MCG
- UPTRAVI TITRATION

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	PULMONARY ARTERIAL HYPERTENSION (PAH): INITIAL: DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	PAH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS
<b>Other Criteria</b>	PAH: INITIAL: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING AGENTS FROM DIFFERENT DRUG CLASSES: 1) FORMULARY VERSION OF AN ORAL ENDOTHELIN RECEPTOR ANTAGONIST, 2) FORMULARY VERSION OF AN ORAL PHOSPHODIESTERASE TYPE-5 INHIBITOR FOR PAH, 3) FORMULARY VERSION OF AN ORAL CGMP STIMULATOR.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# SELINEXOR

## Products Affected

- XPOVIO (100 MG ONCE WEEKLY)  
ORAL TABLET THERAPY PACK 50  
MG
- XPOVIO (40 MG ONCE WEEKLY)  
ORAL TABLET THERAPY PACK 10  
MG, 40 MG
- XPOVIO (40 MG TWICE WEEKLY)  
ORAL TABLET THERAPY PACK 40  
MG
- XPOVIO (60 MG ONCE WEEKLY)  
ORAL TABLET THERAPY PACK 60  
MG
- XPOVIO (60 MG TWICE WEEKLY)
- XPOVIO (80 MG ONCE WEEKLY)  
ORAL TABLET THERAPY PACK 40  
MG
- XPOVIO (80 MG TWICE WEEKLY)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# SELPERCATINIB

## Products Affected

- RETEVMO ORAL CAPSULE 40 MG, 80 MG
- RETEVMO ORAL TABLET 120 MG, 160 MG, 40 MG, 80 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# SELUMETINIB

## Products Affected

- KOSELUGO ORAL CAPSULE 10 MG,  
25 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# SILDENAFIL TABLET

## Products Affected

- *sildenafil citrate oral tablet 20 mg*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	PULMONARY ARTERIAL HYPERTENSION (PAH): INITIAL: AGES 18 YEARS OR OLDER: DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS. AGES 1 TO 17 YEARS: DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PAP GREATER THAN 20 MMHG, 2) PCWP OF 15 MMHG OR LESS, AND 3) PVR OF 3 WOOD UNITS OR GREATER.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	PAH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	PAH: INITIAL/RENEWAL: 1) NOT CONCURRENTLY OR INTERMITTENTLY TAKING ORAL ERECTILE DYSFUNCTION AGENTS (E.G. CIALIS, VIAGRA) OR ANY ORGANIC NITRATES IN ANY FORM AND 2) NO CONCURRENT USE WITH GUANYLATE CYCLASE STIMULATORS.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# SIPONIMOD

## Products Affected

- MAYZENT ORAL TABLET 0.25 MG, 1 MG, 2 MG
- MAYZENT STARTER PACK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# SIROLIMUS PROTEIN-BOUND

---

## Products Affected

- FYARRO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# SODIUM OXYBATE-XYREM

## Products Affected

- *sodium oxybate*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: CATAPLEXY IN NARCOLEPSY, EXCESSIVE DAYTIME SLEEPINESS (EDS) IN NARCOLEPSY: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST, PSYCHIATRIST, OR SPECIALIST IN SLEEP MEDICINE
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
<b>Other Criteria</b>	INITIAL: EDS IN NARCOLEPSY: 1) NO CONCURRENT USE WITH A SEDATIVE HYPNOTIC AGENT, 2) AGES 18 YEARS OR OLDER: TRIAL, FAILURE OF, OR CONTRAINDICATION TO A FORMULARY VERSION OF MODAFINIL, ARMODAFINIL, OR SUNOSI AND ONE GENERIC STIMULANT INDICATED FOR EDS IN NARCOLEPSY, AND 3) AGES 7 TO 17 YEARS: TRIAL, FAILURE OF, OR CONTRAINDICATION TO ONE GENERIC STIMULANT INDICATED FOR EDS IN NARCOLEPSY. CATAPLEXY IN NARCOLEPSY: NO CONCURRENT USE WITH A SEDATIVE HYPNOTIC AGENT. RENEWAL: CATAPLEXY IN NARCOLEPSY, EDS IN NARCOLEPSY: 1) SUSTAINED IMPROVEMENT OF SYMPTOMS COMPARED TO BASELINE, AND 2) NO CONCURRENT USE WITH A SEDATIVE HYPNOTIC AGENT.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# SOFOSBUVIR/VELPATASVIR

## Products Affected

- EPCLUSA ORAL PACKET 150-37.5 MG, 200-50 MG
- EPCLUSA ORAL TABLET

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	HCV RNA LEVEL WITHIN PAST 6 MONTHS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE.
<b>Other Criteria</b>	1) CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE, 2) NOT CONCURRENTLY TAKING ANY OF THE FOLLOWING MEDICATIONS: AMIODARONE, CARBAMAZEPINE, PHENYTOIN, PHENOBARBITAL, OXCARBAZEPINE, RIFAMPIN, RIFABUTIN, RIFAPENTINE, HIV REGIMEN THAT CONTAINS EFAVIRENZ, ROSUVASTATIN AT DOSES ABOVE 10MG, TIPRANA VIR/RITONAVIR, TOPOTECAN, SOVALDI (AS A SINGLE AGENT), HARVONI, ZEPATIER, MAVYRET, OR VOSEVI, AND 3) PATIENTS WITH DECOMPENSATED CIRRHOSIS REQUIRE CONCURRENT RIBAVIRIN UNLESS RIBAVIRIN INELIGIBLE.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# SOFOSBUVIR/VELPATASVIR/VOXILAPREVIR

## Products Affected

- VOSEVI

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	HCV RNA LEVEL WITHIN PAST 6 MONTHS
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE.
<b>Other Criteria</b>	1) CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE, 2) NOT CONCURRENTLY TAKING ANY OF THE FOLLOWING MEDICATIONS: AMIODARONE, CARBAMAZEPINE, PHENYTOIN, PHENOBARBITAL, OXCARBAZEPINE, RIFAMPIN, RIFABUTIN, RIFAPENTINE, CYCLOSPORINE, PITAVASTATIN, PRAVASTATIN (DOSES ABOVE 40MG), ROSUVASTATIN, METHOTREXATE, MITOXANTRONE, IMATINIB, IRINOTECAN, LAPATINIB, SULFASALAZINE, TOPOTECAN, OR HIV REGIMEN THAT CONTAINS EFAVIRENZ, ATAZANAVIR, LOPINAVIR, TIPRANAVIR/RITONAVIR, SOVALDI (AS A SINGLE AGENT), EPCLUSA, HARVONI, ZEPATIER, OR MAVYRET, AND 3) DOES NOT HAVE MODERATE OR SEVERE HEPATIC IMPAIRMENT (CHILD-PUGH B OR C).
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# SOMATROPIN - NORDITROPIN

## Products Affected

- NORDITROPIN FLEXPRO  
SUBCUTANEOUS SOLUTION PEN-  
INJECTOR

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	INITIAL/RENEWAL: ATHLETIC ENHANCEMENT, ANTI-AGING PURPOSES.
<b>Required Medical Information</b>	INITIAL: PEDIATRIC GROWTH HORMONE DEFICIENCY (GHD), IDIOPATHIC SHORT STATURE (ISS), SMALL FOR GESTATIONAL AGE (SGA), TURNER SYNDROME (TS), NOONAN SYNDROME: HEIGHT AT LEAST 2 STANDARD DEVIATIONS BELOW THE MEAN HEIGHT FOR CHILDREN OF THE SAME AGE AND GENDER. PRADER WILLI SYNDROME (PWS): CONFIRMED GENETIC DIAGNOSIS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL/RENEWAL: ALL INDICATIONS: PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: ADULT GHD: GHD ALONE OR ASSOCIATED WITH MULTIPLE HORMONE DEFICIENCIES (HYPOPITUITARISM), AS A RESULT OF PITUITARY DISEASE, HYPOTHALAMIC DISEASE, SURGERY, RADIATION THERAPY, OR TRAUMA, OR FOR CONTINUATION OF THERAPY FROM CHILDHOOD ONSET GHD. PEDIATRIC GHD, ISS, SGA, TS, NOONAN SYNDROME: OPEN EPIPHYSES AS CONFIRMED BY RADIOGRAPH OF THE WRIST AND HAND. RENEWAL: PEDIATRIC GHD: 1) IMPROVEMENT WHILE ON THERAPY (I.E., INCREASED HEIGHT OR INCREASED GROWTH VELOCITY), AND 2) OPEN EPIPHYSES AS CONFIRMED BY RADIOGRAPH OF THE WRIST AND HAND OR HAS NOT COMPLETED PREPUBERTAL GROWTH. ISS, SGA, TS, NOONAN SYNDROME: 1) IMPROVEMENT WHILE ON THERAPY (I.E., INCREASED HEIGHT OR INCREASED GROWTH VELOCITY), AND 2) OPEN EPIPHYSES AS CONFIRMED BY RADIOGRAPH OF

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
	THE WRIST AND HAND. PWS: IMPROVEMENT IN BODY COMPOSITION.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# SOMATROPIN - SEROSTIM

## Products Affected

- SEROSTIM SUBCUTANEOUS SOLUTION RECONSTITUTED 4 MG, 5 MG, 6 MG

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	INITIAL/RENEWAL: ATHLETIC ENHANCEMENT, ANTI-AGING PURPOSES
<b>Required Medical Information</b>	INITIAL: HIV/WASTING: ONE OF THE FOLLOWING FOR WEIGHT LOSS: 1) 10% UNINTENTIONAL WEIGHT LOSS OVER 12 MONTHS, 2) 7.5% UNINTENTIONAL WEIGHT LOSS OVER 6 MONTHS, 3) 5% BODY CELL MASS (BCM) LOSS WITHIN 6 MONTHS, 4) BCM LESS THAN 35% (MEN) OF TOTAL BODY WEIGHT AND BODY MASS INDEX (BMI) LESS THAN 27 KG PER METER SQUARED, 5) BCM LESS THAN 23% (WOMEN) OF TOTAL BODY WEIGHT AND BMI LESS THAN 27 KG PER METER SQUARED, OR 6) BMI LESS THAN 18.5 KG PER METER SQUARED.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	HIV/WASTING: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST, NUTRITIONAL SUPPORT SPECIALIST, OR INFECTIOUS DISEASE SPECIALIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 3 MONTHS.
<b>Other Criteria</b>	HIV/WASTING: INITIAL: 1) INADEQUATE RESPONSE TO ONE PREVIOUS THERAPY (E.G., MEGACE, APPETITE STIMULANTS, ANABOLIC STEROIDS). RENEWAL: 1) CLINICAL BENEFIT IN MUSCLE MASS AND WEIGHT.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# SONIDEGIB

---

## Products Affected

- ODOMZO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	LOCALLY ADVANCED BASAL CELL CARCINOMA (BCC): BASELINE SERUM CREATINE KINASE (CK) AND SERUM CREATININE LEVELS
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# SORAFENIB

## Products Affected

- *sorafenib tosylate*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# SOTATERCEPT-CSRK

## Products Affected

- WINREVAIR

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	PULMONARY ARTERIAL HYPERTENSION (PAH): INITIAL: DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	PAH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	PAH: INITIAL: 1) ON BACKGROUND PAH THERAPY (FOR AT LEAST 3 MONTHS) WITH AT LEAST TWO OF THE FOLLOWING AGENTS FROM DIFFERENT DRUG CLASSES: A) ORAL ENDOTHELIN RECEPTOR ANTAGONIST, B) ORAL PHOSPHODIESTERASE TYPE-5 INHIBITOR FOR PAH, C) ORAL CGMP STIMULATOR, D) IV/SQ PROSTACYCLIN, OR 2) ON ONE AGENT FROM ONE OF THE ABOVE DRUG CLASSES, AND HAS A CONTRAINDICATION OR INTOLERANCE TO ALL OF THE OTHER DRUG CLASSES.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# SOTORASIB

---

## Products Affected

- LUMAKRAS ORAL TABLET 120 MG, 240 MG, 320 MG

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# STIRIPENTOL

## Products Affected

- DIACOMIT ORAL CAPSULE 250 MG, 500 MG
- DIACOMIT ORAL PACKET 250 MG, 500 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	DRAVET SYNDROME: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# SUNITINIB

## Products Affected

- *sunitinib malate*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	GASTROINTESTINAL STROMAL TUMORS (GIST): TRIAL OF OR CONTRAINDICATION TO IMATINIB (GLEEVEC).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TADALAFIL - ADCIRCA, ALYQ

## Products Affected

- *alyq*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	PULMONARY ARTERIAL HYPERTENSION (PAH): INITIAL: DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	PAH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	PAH: INITIAL/RENEWAL: 1) NOT CONCURRENTLY OR INTERMITTENTLY TAKING ORAL ERECTILE DYSFUNCTION AGENTS (E.G. CIALIS, VIAGRA) OR ANY ORGANIC NITRATES IN ANY FORM, AND 2) NO CONCURRENT USE WITH GUANYLATE CYCLASE STIMULATORS.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# TADALAFIL-CIALIS

## Products Affected

- *tadalafil oral tablet 2.5 mg, 5 mg*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	ERECTILE DYSFUNCTION WITHOUT DIAGNOSIS OF BENIGN PROSTATIC HYPERPLASIA (BPH).
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	BPH: 1) TRIAL OF ONE ALPHA BLOCKER (E.G., DOXAZOSIN, TERAZOSIN, TAMSULOSIN, ALFUZOSIN), AND 2) TRIAL OF ONE 5-ALPHA-REDUCTASE INHIBITOR (E.G., FINASTERIDE, DUTASTERIDE). APPLIES TO 2.5MG AND 5MG STRENGTHS ONLY.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# TALAZOPARIB

## Products Affected

- TALZENNA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	ADVANCED OR METASTATIC BREAST CANCER: 1) HAS BEEN TREATED WITH CHEMOTHERAPY IN THE NEOADJUVANT, ADJUVANT, OR METASTATIC SETTING, AND 2) IF HORMONE RECEPTOR (HR)-POSITIVE BREAST CANCER, RECEIVED PRIOR TREATMENT WITH ENDOCRINE THERAPY OR IS CONSIDERED INAPPROPRIATE FOR ENDOCRINE THERAPY. METASTATIC CASTRATION-RESISTANT PROSTATE CANCER: 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# TALQUETAMAB-TGVS

## Products Affected

- TALVEY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TARLATAMAB-DLLE

## Products Affected

- IMDELLTRA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TAZEMETOSTAT

---

## Products Affected

- TAZVERIK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TEBENTAFUSP-TEBN

## Products Affected

- KIMMTRAK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TECLISTAMAB-CQYV

## Products Affected

- TECVAYLI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TELOTRISTAT

---

## Products Affected

- XERMELO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	CARCINOID SYNDROME DIARRHEA: PRESCRIBED BY OR IN CONSULTATION WITH AN ONCOLOGIST OR GASTROENTEROLOGIST
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TEPOTINIB

---

## Products Affected

- TEPMETKO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TESTOSTERONE

## Products Affected

- *testosterone gel 1.62 % transdermal*
- *testosterone transdermal gel 12.5 mg/act (1%), 20.25 mg/act (1.62%), 25 mg/2.5gm (1%), 50 mg/5gm (1%)*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	MALE HYPOGONADISM: INITIAL: CONFIRMED BY: 1) AT LEAST TWO TOTAL SERUM TESTOSTERONE LEVELS OF LESS THAN 300 NG/DL TAKEN ON SEPARATE OCCASIONS, OR 2) FREE SERUM TESTOSTERONE LEVEL OF LESS THAN 5 NG/DL.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS
<b>Other Criteria</b>	MALE HYPOGONADISM: INITIAL: 1) 40 YEARS OR OLDER: PROSTATE SPECIFIC ANTIGEN (PSA) HAS BEEN EVALUATED FOR PROSTATE CANCER SCREENING. RENEWAL: 1) 40 YEARS OR OLDER: PSA HAS BEEN EVALUATED FOR PROSTATE CANCER SCREENING, AND 2) IMPROVED SYMPTOMS COMPARED TO BASELINE AND TOLERANCE TO TREATMENT.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# TESTOSTERONE CYPIONATE

## Products Affected

- *testosterone cypionate intramuscular solution 100 mg/ml, 200 mg/ml, 200 mg/ml (1 ml)*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	MALE HYPOGONADISM: INITIAL: CONFIRMED BY: 1) AT LEAST TWO TOTAL SERUM TESTOSTERONE LEVELS OF LESS THAN 300 NG/DL TAKEN ON SEPARATE OCCASIONS, OR 2) FREE SERUM TESTOSTERONE LEVEL OF LESS THAN 5 NG/DL.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	INITIAL/RENEWAL: 12 MONTHS
<b>Other Criteria</b>	MALE HYPOGONADISM: INITIAL: 1) 40 YEARS OR OLDER: PROSTATE SPECIFIC ANTIGEN (PSA) HAS BEEN EVALUATED FOR PROSTATE CANCER SCREENING. RENEWAL: 1) 40 YEARS OR OLDER: PSA HAS BEEN EVALUATED FOR PROSTATE CANCER SCREENING, AND 2) IMPROVED SYMPTOMS COMPARED TO BASELINE AND TOLERANCE TO TREATMENT.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# TESTOSTERONE ENANTHATE

## Products Affected

- *testosterone enanthate intramuscular solution*
- XYOSTED

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	MALE HYPOGONADISM: INITIAL: CONFIRMED BY: 1) AT LEAST TWO TOTAL SERUM TESTOSTERONE LEVELS OF LESS THAN 300 NG/DL TAKEN ON SEPARATE OCCASIONS, OR 2) FREE SERUM TESTOSTERONE LEVEL OF LESS THAN 5 NG/DL.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	INITIAL/RENEWAL: MALE DELAYED PUBERTY: 6MO, MALE HYPOGONADISM: 12 MO. OTHER INDICATIONS: 12 MO.
<b>Other Criteria</b>	INITIAL: MALE HYPOGONADISM: 1) 40 YEARS OR OLDER: PROSTATE SPECIFIC ANTIGEN (PSA) HAS BEEN EVALUATED FOR PROSTATE CANCER SCREENING. RENEWAL: MALE HYPOGONADISM: 1) 40 YEARS OR OLDER: PSA HAS BEEN EVALUATED FOR PROSTATE CANCER SCREENING, AND 2) IMPROVED SYMPTOMS COMPARED TO BASELINE AND TOLERANCE TO TREATMENT. MALE DELAYED PUBERTY: HAS NOT RECEIVED MORE THAN TWO 6-MONTH COURSES OF TESTOSTERONE REPLACEMENT THERAPY
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# TETRABENAZINE

## Products Affected

- *tetrabenazine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	HUNTINGTONS DISEASE: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST OR MOVEMENT DISORDER SPECIALIST
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# THALIDOMIDE

---

## Products Affected

- THALOMID

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TISLELIZUMAB-JSGR

---

## Products Affected

- TEVIMBRA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TISOTUMAB VEDOTIN-TFTV

## Products Affected

- TIVDAK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TIVOZANIB

---

## Products Affected

- FOTIVDA

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TOCILIZUMAB IV

## Products Affected

- ACTEMRA

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	CORONAVIRUS DISEASE 2019 (COVID-19) IN HOSPITALIZED ADULTS
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: RHEUMATOID ARTHRITIS (RA), POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. SYSTEMIC JUVENILE IDIOPATHIC ARTHRITIS (SJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST, DERMATOLOGIST, OR IMMUNOLOGIST.
<b>Coverage Duration</b>	INITIAL: RA, PJIA, SJIA, GCA: 6 MONTHS. CRS: 1 MONTH. RENEWAL: RA, PJIA, SJIA, GCA: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ, RINVOQ, ORENCIA. PJIA: 1) TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ IR, RINVOQ, ORENCIA, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PJIA. SJIA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR SJIA. RENEWAL: RA: CONTINUES TO BENEFIT FROM THE MEDICATION. PJIA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PJIA. SJIA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
	CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR SJIA.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TOCILIZUMAB SQ

## Products Affected

- ACTEMRA
- ACTEMRA ACTPEN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA), POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. SYSTEMIC JUVENILE IDIOPATHIC ARTHRITIS (SJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST, DERMATOLOGIST, OR IMMUNOLOGIST. SYSTEMIC SCLEROSIS-ASSOCIATED INTERSTITIAL LUNG DISEASE (SSC-ILD): PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR RHEUMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ, RINVOQ, ORENCIA. PJIA: 1) TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ IR, RINVOQ, ORENCIA, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PJIA. SJIA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR SJIA. SSC-ILD: DOES NOT HAVE OTHER KNOWN CAUSES OF INTERSTITIAL LUNG DISEASE (E.G., CONNECTIVE TISSUE DISEASE, DRUG TOXICITY, ASBESTOS OR BERYLLIUM EXPOSURE,

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	HYPERSENSITIVITY PNEUMONITIS). RENEWAL: RA: CONTINUES TO BENEFIT FROM THE MEDICATION. PJIA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PJIA. SJIA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR SJIA. SSC-ILD: CLINICAL MEANINGFUL IMPROVEMENT OR MAINTENANCE IN ANNUAL RATE OF DECLINE.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TOCILIZUMAB-AAZG

## Products Affected

- TYENNE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA), POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. SYSTEMIC JUVENILE IDIOPATHIC ARTHRITIS (SJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST, DERMATOLOGIST, OR IMMUNOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ, RINVOQ, ORENCIA. PJIA: 1) TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ IR, ORENCIA, RINVOQ, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PJIA. SJIA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR SJIA. RENEWAL: RA: CONTINUES TO BENEFIT FROM THE MEDICATION. PJIA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PJIA. SJIA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
	CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR SJIA.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TOCILIZUMAB-AAZG IV

## Products Affected

- TYENNE

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	CORONAVIRUS DISEASE 2019 (COVID-19) IN HOSPITALIZED ADULTS
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: RHEUMATOID ARTHRITIS (RA), POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. SYSTEMIC JUVENILE IDIOPATHIC ARTHRITIS (SJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST, DERMATOLOGIST, OR IMMUNOLOGIST.
<b>Coverage Duration</b>	INITIAL: RA, PJIA, SJIA, GCA: 6 MOS. CRS: 1 MO. RENEWAL: RA, PJIA, SJIA, GCA: 12 MOS.
<b>Other Criteria</b>	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ, RINVOQ, ORENCIA. PJIA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: ENBREL, HUMIRA/CYLTEZO/YUFLYMA, XELJANZ IR, ORENCIA, RINVOQ. CYTOKINE RELEASE SYNDROME (CRS): NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR CRS. INITIAL/RENEWAL FOR PJIA, SJIA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR SAME INDICATION. RENEWAL FOR RA, PJIA, SJIA: CONTINUES TO BENEFIT FROM MEDICATION.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TOFACITINIB

## Products Affected

- XELJANZ
- XELJANZ XR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA), ANKYLOSING SPONDYLITIS (AS), POLYARTICULAR COURSE JUVENILE IDIOPATHIC ARTHRITIS (PCJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST. ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF A PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PSA. PCJIA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PCJIA. AS: 1) TRIAL OF OR CONTRAINDICATION TO AN NSAID, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC. RENEWAL: RA: CONTINUES TO

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	BENEFIT FROM THE MEDICATION. PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. AS: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. PCJIA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PCJIA. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TOPICAL TRETINOIN

## Products Affected

- ALTRENO
- *tretinoin external cream*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	COSMETIC INDICATIONS SUCH AS WRINKLES, PHOTOAGING, MELASMA.
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	ACNE VULGARIS: BRAND TOPICAL TRETINOIN REQUIRES TRIAL OF OR CONTRAINDICATION TO A GENERIC TOPICAL TRETINOIN PRODUCT.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# TORIPALIMAB-TPZI

## Products Affected

- LOQTORZI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	NASOPHARYNGEAL CARCINOMA (NPC): FIRST LINE TREATMENT: 24 MOS, PREVIOUSLY TREATED: LIFETIME.
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TOVORAFENIB

## Products Affected

- OJEMDA ORAL SUSPENSION RECONSTITUTED
- OJEMDA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TRAMADOL

## Products Affected

- TRAMADOL HCL ORAL SOLUTION

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 MONTHS
Other Criteria	PAIN: 1) TRIAL OF OR CONTRAINDICATION TO GENERIC TRAMADOL IMMEDIATE RELEASE TABLET OR GENERIC TRAMADOL/ACETAMINOPHEN COMBINATION PRODUCT, AND 2) UNABLE TO TAKE ORAL SOLID FORMULATIONS OF TRAMADOL OR TRAMADOL/ACETAMINOPHEN COMBINATION PRODUCT (E.G., DIFFICULTY SWALLOWING).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# TRAMETINIB SOLUTION

## Products Affected

- MEKINIST ORAL SOLUTION RECONSTITUTED

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	UNRESECTABLE OR METASTATIC MELANOMA, MELANOMA, METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), LOCALLY ADVANCED OR METASTATIC ANAPLASTIC THYROID CANCER (ATC), UNRESECTABLE OR METASTATIC SOLID TUMOR, LOW-GRADE GLIOMA (LGG): UNABLE TO SWALLOW MEKINIST TABLETS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# TRAMETINIB TABLET

## Products Affected

- MEKINIST ORAL TABLET 0.5 MG, 2 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TRASTUZUMAB-DKST

## Products Affected

- OGIVRI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TRASTUZUMAB-DTTB

## Products Affected

- ONTRUZANT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TRASTUZUMAB-HYALURONIDASE-OYSK

## Products Affected

- HERCEPTIN HYLECTA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	ADJUVANT BREAST CANCER, METASTATIC BREAST CANCER: TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING PREFERRED AGENTS: HERZUMA, OGIVRI, ONTRUZANT, TRAZIMERA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# TRASTUZUMAB-PKRB

## Products Affected

- HERZUMA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TRASTUZUMAB-QYYP

## Products Affected

- TRAZIMERA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TRAZODONE

---

## Products Affected

- RALDESY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	MAJOR DEPRESSIVE DISORDER (MDD); CONTRAINDICATION TO OR UNABLE TO SWALLOW TRAZODONE TABLETS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# TREMELIMUMAB-ACTL

## Products Affected

- IMJUDO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	UHCC: 30 DAYS. METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC): 5 MONTHS.
Other Criteria	UNRESECTABLE HEPATOCELLULAR CARCINOMA (UHCC): HAS NOT RECEIVED PRIOR TREATMENT WITH IMJUDO. NSCLC: HAS NOT RECEIVED A TOTAL OF 5 DOSES OF IMJUDO.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# TRIENTINE CAPSULE

## Products Affected

- *trientine hcl oral capsule 250 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	WILSONS DISEASE: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A HEPATOLOGIST OR GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 12 MONTHS, RENEWAL: LIFETIME.
Other Criteria	WILSONS DISEASE: INITIAL: 1) LEIPZIG SCORE OF 4 OR GREATER, AND 2) TRIAL OF OR CONTRAINDICATION TO FORMULARY VERSION OF PENICILLAMINE TABLET. RENEWAL: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# TRIFLURIDINE/TIPIRACIL

## Products Affected

- LONSURF ORAL TABLET 15-6.14 MG,  
20-8.19 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# TRIPTORELIN-TRELSTAR

## Products Affected

- TRELSTAR MIXJECT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS.
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# TUCATINIB

## Products Affected

- TUKYSA ORAL TABLET 150 MG, 50 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# UBROGEPANT

## Products Affected

- UBRELVY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	ACUTE MIGRAINE TREATMENT: INITIAL: 1) TRIAL OF OR CONTRAINDICATION TO ONE TRIPTAN (E.G., SUMATRIPTAN, RIZATRIPTAN), AND 2) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR ACUTE MIGRAINE TREATMENT. RENEWAL: 1) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR ACUTE MIGRAINE TREATMENT, AND 2) ONE OF THE FOLLOWING: (A) IMPROVEMENT FROM BASELINE IN A VALIDATED ACUTE TREATMENT PATIENT-REPORTED OUTCOME QUESTIONNAIRE, OR (B) THERAPY WORKS CONSISTENTLY IN MAJORITY OF MIGRAINE ATTACKS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# UPADACITINIB

## Products Affected

- RINVOQ
- RINVOQ LQ

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: NON-RADIOGRAPHIC AXIAL SPONDYLOARTHRITIS (NR-AXSPA): 1) C-REACTIVE PROTEIN LEVELS ABOVE THE UPPER LIMIT OF NORMAL, OR 2) SACROILIITIS ON MAGNETIC RESONANCE IMAGING (MRI). ATOPIC DERMATITIS (AD): ATOPIC DERMATITIS COVERING AT LEAST 10 PERCENT OF BODY SURFACE AREA OR ATOPIC DERMATITIS AFFECTING THE FACE, HEAD, NECK, HANDS, FEET, GROIN, OR INTERTRIGINOUS AREAS
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: RHEUMATOID ARTHRITIS (RA), ANKYLOSING SPONDYLITIS (AS), NR-AXSPA, POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS: PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST. AD: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST, ALLERGIST, OR IMMUNOLOGIST. ULCERATIVE COLITIS (UC), CROHNS DISEASE (CD): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF A PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PSA.

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>PJIA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PJIA. AD: 1) INTRACTABLE PRURITUS OR CRACKING/OOZING/BLEEDING OF AFFECTED SKIN, 2) TRIAL OF OR CONTRAINDICATION TO A TOPICAL CORTICOSTEROID, TOPICAL CALCINEURIN INHIBITOR, TOPICAL PDE4 INHIBITOR, OR TOPICAL JAK INHIBITOR, AND 3) NO CONCURRENT USE WITH OTHER SYSTEMIC BIOLOGICS FOR ATOPIC DERMATITIS OR OTHER JAK INHIBITORS FOR ANY INDICATION. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. AS: 1) TRIAL OF OR CONTRAINDICATION TO AN NSAID (NON-STEROIDAL ANTI-INFLAMMATORY DRUG), AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. NR-AXSPA: 1) TRIAL OF OR CONTRAINDICATION TO AN NSAID, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR NR-AXSPA. RENEWAL: RA: CONTINUES TO BENEFIT FROM THE MEDICATION. AD: 1) IMPROVEMENT WHILE ON THERAPY, AND 2) NO CONCURRENT USE WITH OTHER SYSTEMIC BIOLOGICS FOR ATOPIC DERMATITIS OR OTHER JAK INHIBITOR FOR ANY INDICATION. PSA: 1) CONTINUES TO BENEFIT FROM MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. AS: 1) CONTINUES TO BENEFIT MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR AS. NR-AXSPA: 1) CONTINUES TO BENEFIT FROM MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR NR-AXSPA. PJIA: 1) CONTINUES TO BENEFIT FROM MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PJIA. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD.</p>
<b>Indications</b>	All FDA-approved Indications.

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# USTEKINUMAB

## Products Affected

- STELARA SUBCUTANEOUS SOLUTION 45 MG/0.5ML
- STELARA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP OR FACE.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PSA. PSO: 1) ONE OF THE FOLLOWING: (A) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY) FOR THE TREATMENT OF PSO, (B) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA FOR THE TREATMENT OF PSO, OR (C) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	<p>USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC. RENEWAL: PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. PSO: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC.</p>
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# USTEKINUMAB IV

## Products Affected

- STELARA INTRAVENOUS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	2 MONTHS
Other Criteria	CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# USTEKINUMAB-AEKN IV

## Products Affected

- SELARSDI

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP OR FACE.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PSA. PSO: 1) ONE OF THE FOLLOWING: (A) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY) FOR THE TREATMENT OF PSO, (B) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA FOR THE TREATMENT OF PSO, OR (C) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	SMALL MOLECULES FOR UC. RENEWAL: PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. PSO: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# USTEKINUMAB-AEKN SQ

## Products Affected

- SELARSDI

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP OR FACE.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PSA. PSO: 1) ONE OF THE FOLLOWING: (A) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY) FOR THE TREATMENT OF PSO, (B) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA FOR THE TREATMENT OF PSO, OR (C) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	SMALL MOLECULES FOR UC. RENEWAL: PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. PSO: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# USTEKINUMAB-KFCE IV

## Products Affected

- YESINTEK

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP OR FACE.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PSA. PSO: 1) ONE OF THE FOLLOWING: (A) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY) FOR THE TREATMENT OF PSO, (B) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA FOR THE TREATMENT OF PSO, OR (C) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	SMALL MOLECULES FOR UC. RENEWAL: PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. PSO: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# USTEKINUMAB-KFCE SQ

## Products Affected

- YESINTEK

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP OR FACE.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	INITIAL: PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
<b>Coverage Duration</b>	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
<b>Other Criteria</b>	INITIAL: PSA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR PSA. PSO: 1) ONE OF THE FOLLOWING: (A) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY) FOR THE TREATMENT OF PSO, (B) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA FOR THE TREATMENT OF PSO, OR (C) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

PA Criteria	Criteria Details
	SMALL MOLECULES FOR UC. RENEWAL: PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSA. PSO: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR PSO. CD: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR CD. UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR UC.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# VALBENZINE

## Products Affected

- INGREZZA ORAL CAPSULE
- INGREZZA ORAL CAPSULE SPRINKLE
- INGREZZA ORAL CAPSULE THERAPY PACK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	TARDIVE DYSKINESIA (TD): PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST, PSYCHIATRIST, OR MOVEMENT DISORDER SPECIALIST. CHOREA ASSOCIATED WITH HUNTINGTONS DISEASE: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST OR MOVEMENT DISORDER SPECIALIST.
Coverage Duration	12 MONTHS
Other Criteria	TD: HISTORY OF USING AGENTS THAT CAUSE TARDIVE DYSKINESIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# VANDETANIB

---

## Products Affected

- CAPRELSA ORAL TABLET 100 MG,  
300 MG

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	CURRENTLY STABLE ON CAPRELSA REQUIRES NO EXTRA CRITERIA.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# VANZACAFTOR-TEZACAFTOR- DEUTIVACAFTOR (ALYFTREK)

## Products Affected

- ALYFTREK ORAL TABLET 10-50-125  
MG, 4-20-50 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	CF: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR CYSTIC FIBROSIS EXPERT.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: LIFETIME.
Other Criteria	CF: INITIAL: NO CONCURRENT USE WITH ANOTHER CFTR MODULATOR. RENEWAL: 1) IMPROVEMENT IN CLINICAL STATUS, AND 2) NO CONCURRENT USE WITH ANOTHER CFTR MODULATOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# VEMURAFENIB

## Products Affected

- ZELBORAF

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	MELANOMA: ZELBORAF WILL BE USED ALONE OR IN COMBINATION WITH COTELLIC
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# VENETOCLAX

## Products Affected

- VENCLEXTA ORAL TABLET 10 MG, 100 MG, 50 MG
- VENCLEXTA STARTING PACK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# VERICIGUAT

## Products Affected

- VERQUVO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL/RENEWAL:12 MONTHS.
Other Criteria	HEART FAILURE (HF): INITIAL: 1) NO CONCURRENT USE WITH LONG-ACTING NITRATES OR NITRIC OXIDE DONORS, RIOCIQUAT, OR PDE-5 INHIBITORS, 2) TRIAL OF OR CONTRAINDICATION TO ONE PREFERRED SGLT-2 INHIBITOR, AND 3) TRIAL OF OR CONTRAINDICATION TO ONE AGENT FROM ANY OF THE FOLLOWING STANDARD OF CARE CLASSES: (A) ACE INHIBITOR, ARB, OR ARNI, (B) BETA BLOCKER (I.E., BISOPROLOL, CARVEDILOL, METOPROLOL SUCCINATE), OR (C) ALDOSTERONE ANTAGONIST (I.E., SPIRONOLACTONE, EPLERENONE). RENEWAL: NO CONCURRENT USE WITH LONG-ACTING NITRATES OR NITRIC OXIDE DONORS, RIOCIQUAT, OR PDE-5 INHIBITORS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484

Effective: 06/01/2025

Last Updated: 05/07/2025

Y0144\_PACRIT25\_C

# VIGABATRIN

## Products Affected

- *vigabatin*
- *vigadrone*
- *vigpoder*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	REFRACTORY COMPLEX PARTIAL SEIZURES (CPS), INFANTILE SPASMS: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	CPS: TRIAL OF OR CONTRAINDICATION TO TWO ANTIEPILEPTIC AGENTS.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# VIMSELTINIB

---

## Products Affected

- ROMVIMZA

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# VISMODEGIB

---

## Products Affected

- ERIVEDGE

<b>PA Criteria</b>	<b>Criteria Details</b>
<b>Exclusion Criteria</b>	
<b>Required Medical Information</b>	
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 MONTHS
<b>Other Criteria</b>	
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	
<b>Part B Prerequisite</b>	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# VORASIDENIB

---

## Products Affected

- VORANIGO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# VORICONAZOLE SUSPENSION

## Products Affected

- *voriconazole oral suspension reconstituted*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	CANDIDA INFECTIONS: 3 MOS. CONTINUATION OF THERAPY, ALL OTHER INDICATIONS: 6 MOS.
Other Criteria	CANDIDA INFECTIONS: 1) TRIAL OF OR CONTRAINDICATION TO FLUCONAZOLE, AND 2) UNABLE TO SWALLOW TABLETS. ALL INDICATIONS EXCEPT ESOPHAGEAL CANDIDIASIS: UNABLE TO SWALLOW TABLETS. CONTINUATION OF THERAPY AFTER HOSPITAL DISCHARGE REQUIRES NO EXTRA CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
 Effective: 06/01/2025  
 Last Updated: 05/07/2025  
 Y0144\_PACRIT25\_C

# ZANIDATAMAB-HRII

## Products Affected

- ZIIHERA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ZANUBRUTINIB

---

## Products Affected

- BRUKINSA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ZENOCUTUZUMAB-ZBCO

## Products Affected

- BIZENGRI (750 MG DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ZOLBETUXIMAB-CLZB

## Products Affected

- VYLOY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

# ZURANOLONE

---

## Products Affected

- ZURZUVAE ORAL CAPSULE 20 MG, 25 MG, 30 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	14 DAYS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

Formulary IDs: 25482, 25484  
Effective: 06/01/2025  
Last Updated: 05/07/2025  
Y0144\_PACRIT25\_C

## INDEX

### A

abiraterone acetate ..... 6  
abirtega..... 6  
ACTEMRA..... 335, 336, 337, 338  
ACTEMRA ACTPEN ..... 337, 338  
ACTHAR ..... 73  
ACTHAR GEL SUBCUTANEOUS PEN-  
INJECTOR 40 UNIT/0.5ML, 80  
UNIT/ML ..... 73  
ACTIMMUNE..... 175  
ADEMPAS ..... 280, 281  
ADVOCATE INSULIN SYRINGE 29G X  
1/2 ..... 159, 171  
ADVOCATE INSULIN SYRINGE 30G X  
5/16 ..... 159, 171  
ADVOCATE INSULIN SYRINGE 31G X  
5/16 ..... 159, 171  
AIMOVIG..... 119  
AJOVY ..... 135  
AKEEGA ..... 222  
ALCOHOL PREP PAD..... 159, 171  
ALCOHOL PREP PAD 70 %..... 159, 171  
ALCOHOL PREP PADS PAD 70 % .... 159,  
171  
ALCOHOL SWABS PAD..... 159, 171  
ALCOHOL SWABS PAD 70 % .... 159, 171  
ALECENSA..... 20  
ALTRENO..... 345  
ALUNBRIG ORAL TABLET 180 MG, 30  
MG, 90 MG..... 58  
ALUNBRIG ORAL TABLET THERAPY  
PACK..... 58  
ALVAIZ..... 104  
ALYFTREK ORAL TABLET 10-50-125  
MG, 4-20-50 MG ..... 379  
alyq..... 317  
ANKTIVA ..... 228  
AQ INSULIN SYRINGE 31G X 5/16... 159,  
171  
ARCALYST ..... 275, 276  
ARIKAYCE..... 22  
armodafinil..... 214  
ASSURE ID INSULIN SAFETY SYR 29G  
X 1/2..... 159, 171

ASSURE ID INSULIN SAFETY SYR 31G  
X 15/64..... 159, 171  
AUGTYRO ORAL CAPSULE 160 MG, 40  
MG ..... 267  
AUM ALCOHOL PREP PADS PAD 70 %  
..... 159, 171  
AUSTEDO ORAL TABLET 12 MG, 6 MG,  
9 MG ..... 87  
AUSTEDO XR ORAL TABLET  
EXTENDED RELEASE 24 HOUR 12  
MG, 18 MG, 24 MG, 30 MG, 36 MG, 42  
MG, 48 MG, 6 MG ..... 87  
AUSTEDO XR PATIENT TITRATION . 87  
AVONEX PEN INTRAMUSCULAR  
AUTO-INJECTOR KIT..... 172  
AVONEX PREFILLED  
INTRAMUSCULAR PREFILLED  
SYRINGE KIT..... 172  
AYVAKIT ..... 37  
**B**  
BALVERSA ORAL TABLET 3 MG, 4  
MG, 5 MG..... 118  
BD ECLIPSE SYRINGE 30G X 1/2..... 159,  
171  
BD INSULIN SYR ULTRAFINE II 31G X  
5/16 ..... 159, 171  
BD INSULIN SYRINGE 25G X 1. 159, 171  
BD INSULIN SYRINGE 25G X 5/8..... 159,  
171  
BD INSULIN SYRINGE 26G X 1/2..... 159,  
171  
BD INSULIN SYRINGE 27.5G X 5/8.. 159,  
171  
BD INSULIN SYRINGE 27G X 1/2..... 159,  
171  
BD INSULIN SYRINGE 29G X 1/2..... 159,  
171  
BD INSULIN SYRINGE HALF-UNIT 31G  
X 5/16..... 159, 171  
BD INSULIN SYRINGE MICROFINE 27G  
X 5/8..... 159, 171  
BD INSULIN SYRINGE MICROFINE 28G  
X 1/2..... 159, 171

BD INSULIN SYRINGE U-100 1 ML . 159, 171  
 BD INSULIN SYRINGE U-500 .... 159, 171  
 BD INSULIN SYRINGE ULTRAFINE 29G X 1/2..... 159, 160, 171  
 BD INSULIN SYRINGE ULTRAFINE 30G X 1/2..... 160, 171  
 BD INSULIN SYRINGE ULTRAFINE 31G X 5/16..... 160, 171  
 BD SAFETYGLIDE INSULIN SYRINGE 29G X 1/2..... 160, 171  
 BD SAFETYGLIDE INSULIN SYRINGE 30G X 5/16..... 160, 171  
 BD SAFETYGLIDE INSULIN SYRINGE 31G X 15/64..... 160, 171  
 BD SAFETYGLIDE INSULIN SYRINGE 31G X 5/16..... 160, 171  
 BD SAFETYGLIDE SYRINGE/NEEDLE 27G X 5/8..... 160, 171  
 BD SAFETY-LOK INSULIN SYRINGE 29G X 1/2..... 160, 171  
 BD SWAB SINGLE USE REGULAR PAD ..... 160, 171  
 BD SWABS SINGLE USE BUTTERFLY PAD..... 160, 171  
 BD VEO INSULIN SYR U/F 1/2UNIT 31G X 15/64..... 160, 171  
 BD VEO INSULIN SYR ULTRAFINE 31G X 15/64..... 160, 171  
 BD VEO INSULIN SYRINGE U/F 31G X 15/64 ..... 160, 171  
 BENDAMUSTINE HCL INTRAVENOUS SOLUTION..... 46  
 bendamustine hcl intravenous solution reconstituted..... 46  
 BENDEKA ..... 46  
 BENLYSTA SUBCUTANEOUS..... 43  
 BESREMI ..... 290  
 betaine ..... 49  
 BETASERON SUBCUTANEOUS KIT 173  
 bexarotene ..... 53  
 BIZENGRI (750 MG DOSE) ..... 390  
 bortezomib injection ..... 55  
 BORUZU ..... 55  
 bosentan ..... 56

BOSULIF ORAL CAPSULE 100 MG, 50 MG ..... 57  
 BOSULIF ORAL TABLET 100 MG, 400 MG, 500 MG..... 57  
 BRAFTOVI ORAL CAPSULE 75 MG . 108  
 BRUKINSA ..... 389  
**C**  
 CABOMETYX ORAL TABLET 20 MG, 40 MG, 60 MG..... 61  
 CALQUENCE ..... 8  
 CAPRELSA ORAL TABLET 100 MG, 300 MG ..... 378  
 CAREONE INSULIN SYRINGE 30G X 1/2 ..... 160, 171  
 CAREONE INSULIN SYRINGE 31G X 5/16 ..... 160, 171  
 CARETOUCH ALCOHOL PREP PAD 70 % ..... 160, 171  
 CARETOUCH INSULIN SYRINGE 28G X 5/16 ..... 160, 171  
 CARETOUCH INSULIN SYRINGE 29G X 5/16 ..... 160, 171  
 CARETOUCH INSULIN SYRINGE 30G X 5/16 ..... 160, 171  
 CARETOUCH INSULIN SYRINGE 31G X 5/16 ..... 160, 171  
 carglumic acid oral tablet soluble ..... 65  
 CAYSTON..... 41  
 CIMZIA (2 SYRINGE) ..... 67, 68  
 CIMZIA SUBCUTANEOUS KIT 2 X 200 MG ..... 67, 68  
 CINQAIR..... 268, 269  
 COMETRIQ (100 MG DAILY DOSE) ORAL KIT 80 & 20 MG ..... 60  
 COMETRIQ (140 MG DAILY DOSE) ORAL KIT 3 X 20 MG & 80 MG ..... 60  
 COMETRIQ (60 MG DAILY DOSE)..... 60  
 COMFORT ASSIST INSULIN SYRINGE 29G X 1/2..... 160, 171  
 COMFORT ASSIST INSULIN SYRINGE 31G X 5/16..... 160, 171  
 COMFORT EZ INSULIN SYRINGE 28G X 1/2..... 160, 171  
 COMFORT EZ INSULIN SYRINGE 29G X 1/2..... 160, 161, 171

COMFORT EZ INSULIN SYRINGE 30G  
 X 1/2..... 161, 171  
 COMFORT EZ INSULIN SYRINGE 30G  
 X 5/16..... 161, 171  
 COMFORT EZ INSULIN SYRINGE 31G  
 X 15/64..... 161, 171  
 COMFORT EZ INSULIN SYRINGE 31G  
 X 5/16..... 161, 171  
 COPIKTRA..... 99  
 CORTROPHIN ..... 73  
 COSENTYX (300 MG DOSE)..... 296, 297  
 COSENTYX INTRAVENOUS..... 294, 295  
 COSENTYX SENSOREADY (300 MG)  
 ..... 296, 297  
 COSENTYX SUBCUTANEOUS  
 SOLUTION PREFILLED SYRINGE 75  
 MG/0.5ML ..... 296, 297  
 COSENTYX UNOREADY ..... 296, 297  
 COTELLIC ..... 72  
 CURITY ALCOHOL PREPS PAD 70 %  
 ..... 161, 171  
 CURITY ALL PURPOSE SPONGES PAD  
 2..... 161, 171  
 CURITY GAUZE PAD 2 ..... 161, 171  
 CURITY GAUZE SPONGE PAD 2..... 161,  
 171  
 CURITY SPONGES PAD 2..... 161, 171  
 CVS GAUZE PAD 2 ..... 161, 171  
 CVS GAUZE STERILE PAD 2 ..... 161, 171  
 CYLTEZO (2 PEN) ..... 16, 17, 18  
 CYLTEZO (2 SYRINGE) ..... 16, 17, 18  
 CYLTEZO-CD/UC/HS STARTER... 16, 17,  
 18  
 CYLTEZO-PSORIASIS/UV STARTER 16,  
 17, 18  
**D**  
 dalfampridine er ..... 79  
 DANYELZA..... 215  
 DANZITEN ..... 218  
 dasatinib oral tablet 100 mg, 140 mg, 20 mg,  
 50 mg, 70 mg, 80 mg ..... 81  
 DATROWAY ..... 82  
 DAURISMO ORAL TABLET 100 MG, 25  
 MG ..... 142  
 deferasirox granules ..... 84, 85  
 deferasirox oral tablet ..... 84, 85

deferasirox oral tablet soluble 250 mg, 500  
 mg ..... 84, 85  
 DERMACEA GAUZE SPONGE PAD 2  
 ..... 161, 171  
 DERMACEA IV DRAIN SPONGES PAD  
 2..... 161, 171  
 DERMACEA NON-WOVEN SPONGES  
 PAD 2..... 161, 171  
 DERMACEA TYPE VII GAUZE PAD 2  
 ..... 161, 171  
 DIACOMIT ORAL CAPSULE 250 MG,  
 500 MG ..... 315  
 DIACOMIT ORAL PACKET 250 MG, 500  
 MG ..... 315  
 diclofenac epolamine external ..... 89  
 diclofenac sodium external solution 2 % .. 90  
 dimethyl fumarate oral capsule delayed  
 release 120 mg, 240 mg ..... 91  
 dimethyl fumarate starter pack oral capsule  
 delayed release therapy pack ..... 91  
 dronabinol ..... 94  
 DROPLET INSULIN SYRINGE 29G X 1/2  
 ..... 161, 171  
 DROPLET INSULIN SYRINGE 30G X 1/2  
 ..... 161, 171  
 DROPLET INSULIN SYRINGE 30G X  
 15/64 ..... 161, 171  
 DROPLET INSULIN SYRINGE 30G X  
 5/16 ..... 161, 171  
 DROPLET INSULIN SYRINGE 31G X 1/4  
 ..... 161, 171  
 DROPLET INSULIN SYRINGE 31G X  
 15/64 ..... 161, 171  
 DROPLET INSULIN SYRINGE 31G X  
 5/16 ..... 161, 171  
 DROPSAFE ALCOHOL PREP PAD 70 %  
 ..... 161, 171  
 DROPSAFE SAFETY SYRINGE/NEEDLE  
 29G X 1/2..... 162, 171  
 DROPSAFE SAFETY SYRINGE/NEEDLE  
 31G X 15/64..... 162, 171  
 DROPSAFE SAFETY SYRINGE/NEEDLE  
 31G X 5/16..... 162, 171  
 droxidopa ..... 95  
 DRUG MART ULTRA COMFORT SYR  
 29G X 1/2..... 162, 171

DRUG MART ULTRA COMFORT SYR  
 30G X 5/16..... 162, 171  
 DUPIXENT SUBCUTANEOUS  
 SOLUTION AUTO-INJECTOR .... 96, 98  
 DUPIXENT SUBCUTANEOUS  
 SOLUTION PREFILLED SYRINGE . 96,  
 98

**E**

EASY COMFORT ALCOHOL PADS PAD  
 ..... 162, 171  
 EASY COMFORT INSULIN SYRINGE  
 29G X 5/16..... 162, 171  
 EASY COMFORT INSULIN SYRINGE  
 30G X 1/2..... 162, 171  
 EASY COMFORT INSULIN SYRINGE  
 30G X 5/16..... 162, 171  
 EASY COMFORT INSULIN SYRINGE  
 31G X 1/2..... 162, 171  
 EASY COMFORT INSULIN SYRINGE  
 31G X 5/16..... 162, 171  
 EASY COMFORT INSULIN SYRINGE  
 32G X 5/16..... 162, 171  
 EASY TOUCH ALCOHOL PREP  
 MEDIUM PAD 70 %..... 162, 171  
 EASY TOUCH FLIPLOCK INSULIN SY  
 29G X 1/2..... 162, 171  
 EASY TOUCH FLIPLOCK INSULIN SY  
 30G X 1/2..... 162, 171  
 EASY TOUCH FLIPLOCK INSULIN SY  
 30G X 5/16..... 162, 171  
 EASY TOUCH FLIPLOCK INSULIN SY  
 31G X 5/16..... 162, 171  
 EASY TOUCH FLIPLOCK SAFETY SYR  
 27G X 1/2..... 162, 171  
 EASY TOUCH INSULIN BARRELS U-  
 100 1 ML..... 162, 171  
 EASY TOUCH INSULIN SAFETY SYR  
 29G X 1/2..... 162, 171  
 EASY TOUCH INSULIN SAFETY SYR  
 30G X 1/2..... 162, 171  
 EASY TOUCH INSULIN SAFETY SYR  
 30G X 5/16..... 162, 171  
 EASY TOUCH INSULIN SYRINGE 27G  
 X 1/2..... 162, 171  
 EASY TOUCH INSULIN SYRINGE 27G  
 X 5/8..... 162, 171

EASY TOUCH INSULIN SYRINGE 28G  
 X 1/2..... 162, 171  
 EASY TOUCH INSULIN SYRINGE 29G  
 X 1/2..... 162, 171  
 EASY TOUCH INSULIN SYRINGE 30G  
 X 1/2..... 162, 163, 171  
 EASY TOUCH INSULIN SYRINGE 30G  
 X 5/16..... 163, 171  
 EASY TOUCH INSULIN SYRINGE 31G  
 X 5/16..... 163, 171  
 EASY TOUCH SHEATHLOCK SYRINGE  
 29G X 1/2..... 163, 171  
 EASY TOUCH SHEATHLOCK SYRINGE  
 30G X 1/2..... 163, 171  
 EASY TOUCH SHEATHLOCK SYRINGE  
 30G X 5/16..... 163, 171  
 EASY TOUCH SHEATHLOCK SYRINGE  
 31G X 5/16..... 163, 171  
 ELAHERE ..... 211  
 ELIGARD..... 190  
 ELREXFIO SUBCUTANEOUS  
 SOLUTION 44 MG/1.1ML, 76  
 MG/1.9ML..... 103  
 EMBECTA INS SYR U/F 1/2 UNIT 31G X  
 15/64 ..... 163, 171  
 EMBECTA INS SYR U/F 1/2 UNIT 31G X  
 5/16 ..... 163, 171  
 EMBECTA INSULIN SYR ULTRAFINE  
 30G X 1/2..... 163, 171  
 EMBECTA INSULIN SYR ULTRAFINE  
 31G X 5/16..... 163, 171  
 EMBECTA INSULIN SYRINGE 28G X  
 1/2 ..... 163, 171  
 EMBECTA INSULIN SYRINGE U-100  
 27G X 5/8..... 163, 171  
 EMBECTA INSULIN SYRINGE U-100  
 28G X 1/2..... 163, 171  
 EMGALITY ..... 138  
 EMGALITY (300 MG DOSE) ..... 138  
 ENBREL MINI..... 122, 123  
 ENBREL SUBCUTANEOUS SOLUTION  
 25 MG/0.5ML..... 122, 123  
 ENBREL SUBCUTANEOUS SOLUTION  
 PREFILLED SYRINGE ..... 122, 123  
 ENBREL SUBCUTANEOUS SOLUTION  
 RECONSTITUTED ..... 122, 123

ENBREL SURECLICK SUBCUTANEOUS SOLUTION AUTO-INJECTOR 122, 123  
 EPCLUSA ORAL PACKET 150-37.5 MG, 200-50 MG..... 306  
 EPCLUSA ORAL TABLET..... 306  
 EPIDIOLEX..... 62  
 EPKINLY ..... 113  
 EQL ALCOHOL SWABS PAD 70 %... 163, 171  
 EQL GAUZE PAD 2 ..... 163, 171  
 EQL INSULIN SYRINGE 30G X 5/16. 163, 171  
 ERBITUX ..... 69  
 ERIVEDGE..... 385  
 ERLEADA ORAL TABLET 240 MG, 60 MG ..... 26  
 erlotinib hcl oral tablet 100 mg, 150 mg, 25 mg ..... 120  
 everolimus oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg ..... 124  
 everolimus oral tablet soluble ..... 125  
 EVRYSDI ORAL SOLUTION RECONSTITUTED ..... 285

**F**  
 FASENRA ..... 47, 48  
 FASENRA PEN..... 47, 48  
 fentanyl citrate buccal lozenge on a handle ..... 129  
 fingolimod hcl..... 133  
 FINTEPLA..... 128  
 FOTIVDA ..... 334  
 FREESTYLE PRECISION INS SYR 30G X 5/16 ..... 163, 171  
 FREESTYLE PRECISION INS SYR 31G X 5/16 ..... 163, 171  
 FRUZAQLA ORAL CAPSULE 1 MG, 5 MG ..... 136  
 FYARRO ..... 304

**G**  
 GAUZE PADS PAD 2..... 163, 171  
 GAUZE TYPE VII MEDI-PAK PAD 2 163, 171  
 GAVRETO ..... 261  
 gefitinib ..... 140  
 GILOTRIF ..... 19

glatiramer acetate subcutaneous solution prefilled syringe 20 mg/ml, 40 mg/ml 143  
 glatopa subcutaneous solution prefilled syringe 20 mg/ml, 40 mg/ml..... 143  
 GLOBAL ALCOHOL PREP EASE163, 171  
 GLOBAL EASY GLIDE INSULIN SYR 31G X 15/64..... 163, 171  
 GLOBAL INJECT EASE INSULIN SYR 28G X 1/2..... 163, 171  
 GLOBAL INJECT EASE INSULIN SYR 29G X 1/2..... 163, 171  
 GLOBAL INJECT EASE INSULIN SYR 30G X 5/16..... 163, 171  
 GLUCOPRO INSULIN SYRINGE 30G X 1/2 ..... 163, 171  
 GLUCOPRO INSULIN SYRINGE 30G X 5/16 ..... 164, 171  
 GLUCOPRO INSULIN SYRINGE 31G X 5/16 ..... 164, 171  
 GNP ALCOHOL SWABS PAD..... 164, 171  
 GNP INSULIN SYRINGE 28G X 1/2 .. 164, 171  
 GNP INSULIN SYRINGE 29G X 1/2 .. 164, 171  
 GNP INSULIN SYRINGE 30G X 5/16 164, 171  
 GNP INSULIN SYRINGES 29GX1/2.. 164, 171  
 GNP INSULIN SYRINGES 30G X 5/16 ..... 164, 171  
 GNP INSULIN SYRINGES 30GX5/16 164, 171  
 GNP INSULIN SYRINGES 31GX5/16 164, 171  
 GNP STERILE GAUZE PAD 2..... 164, 171  
 GNP ULTRA COM INSULIN SYRINGE 29G X 1/2..... 164, 171  
 GNP ULTRA COM INSULIN SYRINGE 30G X 5/16..... 164, 171  
 GOMEKLI ORAL CAPSULE 1 MG, 2 MG ..... 210  
 GOMEKLI ORAL TABLET SOLUBLE210  
 GOODSENSE ALCOHOL SWABS PAD 70 % ..... 164, 171

<b>H</b>	
HAEGARDA SUBCUTANEOUS SOLUTION RECONSTITUTED 2000 UNIT, 3000 UNIT .....	59
HARVONI ORAL PACKET 33.75-150 MG, 45-200 MG .....	184
HARVONI ORAL TABLET .....	184
HEALTHWISE INSULIN SYR/NEEDLE 30G X 5/16.....	164, 171
HEALTHWISE INSULIN SYR/NEEDLE 31G X 5/16.....	164, 171
H-E-B INCONTROL ALCOHOL PAD	164, 171
HERCEPTIN HYLECTA.....	353
HERZUMA.....	354
HM STERILE PADS PAD 2.....	164, 171
HM ULTICARE INSULIN SYRINGE 30G X 1/2.....	164, 171
HM ULTICARE INSULIN SYRINGE 31G X 5/16.....	164, 171
HUMIRA (2 PEN) SUBCUTANEOUS AUTO-INJECTOR KIT.....	10, 12
HUMIRA (2 SYRINGE) SUBCUTANEOUS PREFILLED SYRINGE KIT 10 MG/0.1ML, 20 MG/0.2ML, 40 MG/0.4ML, 40 MG/0.8ML .....	10, 12
HUMIRA-CD/UC/HS STARTER SUBCUTANEOUS AUTO-INJECTOR KIT .....	10, 12
HUMIRA-PED<40KG CROHNS STARTER.....	10, 12
HUMIRA-PED>/=40KG CROHNS START .....	10, 12
HUMIRA-PED>/=40KG UC STARTER SUBCUTANEOUS AUTO-INJECTOR KIT .....	10, 12
HUMIRA-PS/UV/ADOL HS STARTER SUBCUTANEOUS AUTO-INJECTOR KIT .....	10, 12
HUMIRA-PSORIASIS/UEVEIT STARTER SUBCUTANEOUS AUTO-INJECTOR KIT .....	10, 12
<b>I</b>	
IBRANCE.....	242
icatibant acetate.....	151

ICLUSIG.....	259
IDHIFA .....	107
imatinib mesylate oral tablet 100 mg, 400 mg .....	153
IMBRUVICA ORAL CAPSULE 140 MG, 70 MG .....	150
IMBRUVICA ORAL SUSPENSION....	150
IMBRUVICA ORAL TABLET .....	150
IMDELLTRA .....	321
IMJUDO .....	357
IMKELDI.....	154
IMPAVIDO.....	209
INCRELEX.....	203
infliximab.....	157, 158
INGREZZA ORAL CAPSULE.....	377
INGREZZA ORAL CAPSULE SPRINKLE .....	377
INGREZZA ORAL CAPSULE THERAPY PACK.....	377
INLYTA ORAL TABLET 1 MG, 5 MG..	39
INQOVI .....	83
INREBIC.....	127
INSULIN SYRINGE 29G X 1/2 ....	164, 171
INSULIN SYRINGE 30G X 5/16 ..	164, 171
INSULIN SYRINGE 31G X 5/16 ..	164, 171
INSULIN SYRINGE/NEEDLE 27G X 1/2 .....	164, 171
INSULIN SYRINGE/NEEDLE 28G X 1/2 .....	165, 171
INSULIN SYRINGE-NEEDLE U-100 27G X 1/2.....	164, 171
INSULIN SYRINGE-NEEDLE U-100 28G X 1/2.....	164, 171
INSULIN SYRINGE-NEEDLE U-100 30G X 5/16.....	164, 171
INSULIN SYRINGE-NEEDLE U-100 31G X 1/4.....	164, 171
INSULIN SYRINGE-NEEDLE U-100 31G X 5/16.....	164, 171
ITOVEBI ORAL TABLET 3 MG, 9 MG .....	156
IWILFIN .....	100
<b>J</b>	
J & J GAUZE PAD 2.....	165, 171
JAKAFI.....	292
javygtor oral tablet .....	293

JAYPIRCA ORAL TABLET 100 MG, 50  
 MG ..... 257  
 JEMPERLI ..... 93  
**K**  
 KALYDECO ..... 177  
 KENDALL HYDROPHILIC FOAM  
 DRESS PAD 2 ..... 165, 171  
 KENDALL HYDROPHILIC FOAM PLUS  
 PAD 2 ..... 165, 171  
 KERENDIA ..... 132  
 KESIMPTA ..... 231  
 KEYTRUDA INTRAVENOUS  
 SOLUTION ..... 250  
 KIMMTRAK ..... 323  
 KINERET SUBCUTANEOUS SOLUTION  
 PREFILLED SYRINGE ..... 24, 25  
 KINRAY INSULIN SYRINGE 29G X 1/2  
 ..... 165, 171  
 KISQALI (200 MG DOSE) ..... 272  
 KISQALI (400 MG DOSE) ..... 272  
 KISQALI (600 MG DOSE) ..... 272  
 KISQALI FEMARA (200 MG DOSE) .. 273  
 KISQALI FEMARA (400 MG DOSE) .. 273  
 KISQALI FEMARA (600 MG DOSE) .. 273  
 KMART VALU INSULIN SYRINGE 29G  
 U-100 1 ML ..... 165, 171  
 KMART VALU INSULIN SYRINGE 30G  
 U-100 0.3 ML ..... 165, 171  
 KMART VALU INSULIN SYRINGE 30G  
 U-100 1 ML ..... 165, 171  
 KOSELUGO ORAL CAPSULE 10 MG, 25  
 MG ..... 301  
 KRAZATI ..... 9  
 KYNMOBI ..... 28  
 KYNMOBI TITRATION KIT ..... 28  
**L**  
 LANREOTIDE ACETATE ..... 180  
 lapatinib ditosylate ..... 181  
 LAZCLUZE ORAL TABLET 240 MG, 80  
 MG ..... 183  
 lenalidomide ..... 185  
 LENVIMA (10 MG DAILY DOSE) ..... 186  
 LENVIMA (12 MG DAILY DOSE) ..... 186  
 LENVIMA (14 MG DAILY DOSE) ..... 186  
 LENVIMA (18 MG DAILY DOSE) ..... 186  
 LENVIMA (20 MG DAILY DOSE) ..... 186

LENVIMA (24 MG DAILY DOSE) ..... 186  
 LENVIMA (4 MG DAILY DOSE) ..... 186  
 LENVIMA (8 MG DAILY DOSE) ..... 186  
 LEUPROLIDE ACETATE (3 MONTH) 189  
 leuprolide acetate injection ..... 188  
 l-glutamine oral packet ..... 195  
 LITETOUCH INSULIN SYRINGE 28G X  
 1/2 ..... 165, 171  
 LITETOUCH INSULIN SYRINGE 29G X  
 1/2 ..... 165, 171  
 LITETOUCH INSULIN SYRINGE 30G X  
 5/16 ..... 165, 171  
 LITETOUCH INSULIN SYRINGE 31G X  
 5/16 ..... 165, 171  
 LIVTENCITY ..... 202  
 LONSURF ORAL TABLET 15-6.14 MG,  
 20-8.19 MG ..... 359  
 LOQTORZI ..... 346  
 LORBRENA ORAL TABLET 100 MG, 25  
 MG ..... 197  
 LUMAKRAS ORAL TABLET 120 MG,  
 240 MG, 320 MG ..... 314  
 LUNSUMIO ..... 213  
 LUPRON DEPOT (1-MONTH) ..... 191, 192  
 LUPRON DEPOT (3-MONTH) ..... 191, 192  
 LUPRON DEPOT (4-MONTH) ..... 191, 192  
 LUPRON DEPOT (6-MONTH) ..... 191, 192  
 LUPRON DEPOT-PED (3-MONTH) ... 193,  
 194  
 LUPRON DEPOT-PED (6-MONTH) ... 193,  
 194  
 LUTRATE DEPOT ..... 189  
 LYBALVI ..... 232  
 LYNPARZA ORAL TABLET ..... 233  
 LYTGOBI (12 MG DAILY DOSE) ..... 137  
 LYTGOBI (16 MG DAILY DOSE) ..... 137  
 LYTGOBI (20 MG DAILY DOSE) ..... 137  
**M**  
 MAGELLAN INSULIN SAFETY SYR  
 29G X 1/2 ..... 165, 171  
 MAGELLAN INSULIN SAFETY SYR  
 30G X 5/16 ..... 165, 171  
 MARGENZA ..... 201  
 MAVENCLAD (10 TABS) ..... 70  
 MAVENCLAD (4 TABS) ..... 70  
 MAVENCLAD (5 TABS) ..... 70

MAVENCLAD (6 TABS) .....	70	MOUNJARO SUBCUTANEOUS	
MAVENCLAD (7 TABS) .....	70	SOLUTION AUTO-INJECTOR .....	146
MAVENCLAD (8 TABS) .....	70	MVASI.....	51
MAVENCLAD (9 TABS) .....	70	<b>N</b>	
MAXI-COMFORT INSULIN SYRINGE		NATPARA.....	243
28G X 1/2.....	165, 171	NERLYNX .....	216
MAXICOMFORT SYR 27G X 1/2	165, 171	NEULASTA ONPRO.....	247
MAYZENT ORAL TABLET 0.25 MG, 1		NIKTIMVO .....	38
MG, 2 MG.....	303	NINLARO.....	179
MAYZENT STARTER PACK.....	303	nitisinone.....	224
MEDIC INSULIN SYRINGE 30G X 5/16		NIVESTYM.....	131
.....	165, 171	NORDITROPIN FLEXPRO	
MEDPURA ALCOHOL PADS 70 %		SUBCUTANEOUS SOLUTION PEN-	
EXTERNAL .....	165, 171	INJECTOR.....	308, 309
MEIJER ALCOHOL SWABS PAD 70 %		NUBEQA.....	80
.....	165, 171	NUCALA SUBCUTANEOUS SOLUTION	
MEKINIST ORAL SOLUTION		AUTO-INJECTOR .....	205, 206
RECONSTITUTED.....	349	NUCALA SUBCUTANEOUS SOLUTION	
MEKINIST ORAL TABLET 0.5 MG, 2		PREFILLED SYRINGE 100 MG/ML, 40	
MG .....	350	MG/0.4ML.....	205, 206
MEKTOVI .....	54	NUCALA SUBCUTANEOUS SOLUTION	
mifepristone oral tablet 300 mg .....	208	RECONSTITUTED .....	205, 206
MIPLYFFA.....	31	NUDEXTA.....	88
MIRASORB SPONGES 2.....	165, 171	NUPLAZID ORAL CAPSULE.....	255
modafinil oral tablet 100 mg, 200 mg.....	214	NUPLAZID ORAL TABLET 10 MG... ..	255
MONOJECT INSULIN SYRINGE 25G X		NURTEC.....	278, 279
5/8 .....	165, 171	NYVEPRIA .....	246
MONOJECT INSULIN SYRINGE 27G X		<b>O</b>	
1/2 .....	165, 171	OCREVUS.....	229
MONOJECT INSULIN SYRINGE 28G X		OCREVUS ZUNOVO .....	230
1/2 .....	165, 171	ODOMZO.....	311
MONOJECT INSULIN SYRINGE 29G X		OFEV .....	219, 220
1/2 .....	165, 171	OGIVRI.....	351
MONOJECT INSULIN SYRINGE 30G X		OGSIVEO ORAL TABLET 100 MG, 150	
5/16 .....	165, 166, 171	MG, 50 MG.....	223
MONOJECT INSULIN SYRINGE 31G X		OJEMDA ORAL SUSPENSION	
5/16 .....	166, 171	RECONSTITUTED .....	347
MONOJECT INSULIN SYRINGE U-100 1		OJEMDA ORAL TABLET .....	347
ML.....	166, 171	OJJAARA .....	212
MONOJECT ULTRA COMFORT		ONAPGO.....	27
SYRINGE 28G X 1/2 .....	166, 171	ONTRUZANT .....	352
MONOJECT ULTRA COMFORT		ONUREG.....	40
SYRINGE 29G X 1/2 .....	166, 171	OPDIVO .....	225
MONOJECT ULTRA COMFORT		OPDIVO QVANTIG .....	226
SYRINGE 30G X 5/16 .....	166, 171	OPDUALAG.....	227
		OPSUMIT .....	200

ORENCIA CLICKJECT.....	3, 4	PRECISION SURE-DOSE SYRINGE 29G X 1/2.....	166, 171
ORENCIA INTRAVENOUS .....	1, 2	PRECISION SURE-DOSE SYRINGE 30G X 3/8.....	166, 171
ORENCIA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE .....	3, 4	PRECISION SURE-DOSE SYRINGE 30G X 5/16.....	166, 171
ORFADIN ORAL SUSPENSION.....	224	PREFERRED PLUS INSULIN SYRINGE 28G X 1/2.....	166, 171
ORGOVYX.....	266	PREVYMIS ORAL TABLET .....	187
ORLISSA ORAL TABLET 150 MG, 200 MG .....	102	PRO COMFORT ALCOHOL PAD 70 % .....	166, 171
ORKAMBI ORAL TABLET .....	199	PRO COMFORT INSULIN SYRINGE 30G X 1/2.....	166, 171
ORSERDU ORAL TABLET 345 MG, 86 MG .....	101	PRO COMFORT INSULIN SYRINGE 30G X 5/16.....	166, 171
OTEZLA .....	29, 30	PRO COMFORT INSULIN SYRINGE 31G X 5/16.....	166, 171
oxandrolone oral .....	240	PROCRIT INJECTION SOLUTION 10000 UNIT/ML, 2000 UNIT/ML, 20000 UNIT/ML, 3000 UNIT/ML, 4000 UNIT/ML, 40000 UNIT/ML .....	114, 115
OZEMPIC (0.25 OR 0.5 MG/DOSE).....	145	PRODIGY INSULIN SYRINGE 28G X 1/2 .....	166, 171
OZEMPIC (1 MG/DOSE) .....	145	PRODIGY INSULIN SYRINGE 31G X 5/16 .....	166, 171
OZEMPIC (2 MG/DOSE) .....	145	PROMACTA ORAL PACKET 12.5 MG, 25 MG .....	105, 106
<b>P</b>		PROMACTA ORAL TABLET 12.5 MG, 25 MG, 50 MG, 75 MG .....	105, 106
pazopanib hcl .....	245	PURE COMFORT ALCOHOL PREP PAD .....	166, 171
PEGASYS SUBCUTANEOUS SOLUTION 180 MCG/ML .....	248	pyrimethamine oral .....	262
PEGASYS SUBCUTANEOUS SOLUTION PREFILLED SYRINGE .....	248	<b>Q</b>	
PEMAZYRE .....	251	QC ALCOHOL.....	166, 171
penicillamine oral tablet.....	252, 253	QC ALCOHOL SWABS PAD 70 % .....	166, 171
PIQRAY (200 MG DAILY DOSE).....	21	QC BORDER ISLAND GAUZE PAD 2 .....	166, 171
PIQRAY (250 MG DAILY DOSE).....	21	QINLOCK.....	282
PIQRAY (300 MG DAILY DOSE).....	21	quinine sulfate oral.....	263
pirfenidone oral capsule .....	256	QULIPTA .....	35
pirfenidone oral tablet 267 mg, 534 mg, 801 mg .....	256	<b>R</b>	
PLEGRIDY STARTER PACK SUBCUTANEOUS SOLUTION AUTO- INJECTOR.....	174	RA ALCOHOL SWABS PAD 70 % .....	166, 171
PLEGRIDY STARTER PACK SUBCUTANEOUS SOLUTION PREFILLED SYRINGE .....	174	RA INSULIN SYRINGE 29G X 1/2.....	166, 171
PLEGRIDY SUBCUTANEOUS SOLUTION AUTO-INJECTOR .....	174		
PLEGRIDY SUBCUTANEOUS SOLUTION PREFILLED SYRINGE	174		
POMALYST .....	258		
posaconazole oral tablet delayed release	260		
PRECISION SUREDOSE PLUS SYR 29G X 1/2.....	166, 171		
PRECISION SURE-DOSE SYRINGE 28G X 1/2.....	166, 171		

RA INSULIN SYRINGE 30G X 5/16... 166, 171	RYBELSUS (FORMULATION R2)..... 145
ra isopropyl alcohol wipes ..... 166, 171	RYBREVANT ..... 23
RA STERILE PAD 2 ..... 166, 171	RYDAPT..... 207
RALDESY ..... 356	RYTELO..... 155
REALITY INSULIN SYRINGE 28G X 1/2 ..... 166, 171	<b>S</b>
REALITY INSULIN SYRINGE 29G X 1/2 ..... 166, 171	SAFETY INSULIN SYRINGES 29G X 1/2 ..... 167, 171
REALITY SWABS PAD..... 166, 171	SAFETY INSULIN SYRINGES 30G X 1/2 ..... 167, 171
RELION ALCOHOL SWABS PAD..... 167, 171	SAFETY INSULIN SYRINGES 30G X 5/16 ..... 167, 171
RELI-ON INSULIN SYRINGE 29G 0.3 ML..... 166, 171	sapropterin dihydrochloride oral tablet... 293
RELI-ON INSULIN SYRINGE 29G 0.5 ML..... 166, 171	SB ALCOHOL PREP PAD 70 %... 167, 171
RELI-ON INSULIN SYRINGE 29G X 1/2 ..... 166, 171	SB INSULIN SYRINGE 29G X 1/2 ..... 167, 171
RELION INSULIN SYRINGE 31G X 15/64 ..... 167, 171	SB INSULIN SYRINGE 30G X 5/16 ... 167, 171
RESTORE CONTACT LAYER PAD 2 167, 171	SB INSULIN SYRINGE 31G X 5/16 ... 167, 171
RETACRIT INJECTION SOLUTION	SCSEMBLIX ORAL TABLET 100 MG, 20 MG, 40 MG..... 32
10000 UNIT/ML, 10000	SECURESAFE INSULIN SYRINGE 29G X 1/2..... 167, 171
UNIT/ML(1ML), 2000 UNIT/ML, 20000	SELARSDI ..... 369, 370, 371, 372
UNIT/ML, 3000 UNIT/ML, 4000	SEROSTIM SUBCUTANEOUS SOLUTION RECONSTITUTED 4 MG, 5 MG, 6 MG..... 310
UNIT/ML, 40000 UNIT/ML ..... 116, 117	SIGNIFOR ..... 244
RETEVMO ORAL CAPSULE 40 MG, 80 MG ..... 300	sildenafil citrate oral tablet 20 mg ..... 302
RETEVMO ORAL TABLET 120 MG, 160 MG, 40 MG, 80 MG ..... 300	SIRTURO ..... 42
REVUFORJ ORAL TABLET 110 MG, 160 MG, 25 MG..... 271	SKYRIZI..... 283, 284
REZLIDHIA ..... 234	SKYRIZI (150 MG DOSE) ..... 283, 284
REZUROCK..... 44	SKYRIZI PEN ..... 283, 284
RIABNI..... 288	SM ALCOHOL PREP PAD ..... 167, 171
RINVOQ..... 363, 364, 365	SM ALCOHOL PREP PAD 6-70 % EXTERNAL ..... 167, 171
RINVOQ LQ..... 363, 364, 365	SM ALCOHOL PREP PAD 70 %.. 167, 171
RITUXAN HYCELA ..... 286	SM GAUZE PAD 2 ..... 167, 171
ROMVIMZA ..... 384	sodium oxybate ..... 305
ROZLYTREK ORAL CAPSULE 100 MG, 200 MG ..... 109	SOMATULINE DEPOT SUBCUTANEOUS SOLUTION 60 MG/0.2ML, 90 MG/0.3ML ..... 180
ROZLYTREK ORAL PACKET ..... 110	SOMAVERT..... 249
RUBRACA ..... 291	sorafenib tosylate ..... 312
RUXIENCE ..... 289	SPRAVATO (56 MG DOSE)..... 121
RYBELSUS ..... 145	SPRAVATO (84 MG DOSE)..... 121

STELARA INTRAVENOUS .....	368	TECHLITE INSULIN SYRINGE 29G X	
STELARA SUBCUTANEOUS		1/2 .....	167, 171
SOLUTION 45 MG/0.5ML .....	366, 367	TECVAYLI.....	324
STELARA SUBCUTANEOUS		TEPMETKO .....	326
SOLUTION PREFILLED SYRINGE	366,	TERUMO INSULIN SYRINGE 29G X 1/2	
367		.....	168, 171
STERILE GAUZE PAD 2 .....	167, 171	testosterone cypionate intramuscular	
STERILE PAD 2.....	167, 171	solution 100 mg/ml, 200 mg/ml, 200	
STIVARGA .....	265	mg/ml (1 ml) .....	328
STRENSIQ .....	33, 34	testosterone enanthate intramuscular	
sunitinib malate.....	316	solution.....	329
SURE COMFORT ALCOHOL PREP PAD		testosterone gel 1.62 % transdermal .....	327
70 % .....	167, 171	testosterone transdermal gel 12.5 mg/act	
SURE COMFORT INSULIN SYRINGE		(1%), 20.25 mg/act (1.62%), 25	
28G X 1/2.....	167, 171	mg/2.5gm (1%), 50 mg/5gm (1%).....	327
SURE COMFORT INSULIN SYRINGE		tetrabenazine .....	330
29G X 1/2.....	167, 171	TEVIMBRA.....	332
SURE COMFORT INSULIN SYRINGE		THALOMID .....	331
30G X 1/2.....	167, 171	THERAGAUZE PAD 2.....	168, 171
SURE COMFORT INSULIN SYRINGE		TIBSOVO .....	178
30G X 5/16.....	167, 171	TIGLUTIK .....	277
SURE COMFORT INSULIN SYRINGE		TIVDAK .....	333
31G X 1/4.....	167, 171	TOPCARE ULTRA COMFORT INS SYR	
SURE COMFORT INSULIN SYRINGE		29G X 1/2.....	168, 171
31G X 5/16.....	167, 171	TOPCARE ULTRA COMFORT INS SYR	
SURE-JECT INSULIN SYRINGE 31G X		30G X 5/16.....	168, 171
5/16 .....	167, 171	TOPCARE ULTRA COMFORT INS SYR	
SURE-PREP ALCOHOL PREP PAD 70 %		31G X 5/16.....	168, 171
.....	167, 171	torpenz oral tablet 10 mg, 2.5 mg, 5 mg, 7.5	
SURGICAL GAUZE SPONGE PAD 2	167,	mg .....	124
171		TRAMADOL HCL ORAL SOLUTION	348
SYMPAZAN.....	71	TRAZIMERA .....	355
SYNRIBO .....	235	TRELSTAR MIXJECT .....	360
<b>T</b>		TREMFYA INTRAVENOUS .....	148, 149
TABRECTA .....	64	TREMFYA ONE-PRESS.....	148, 149
tadalafil oral tablet 2.5 mg, 5 mg .....	318	TREMFYA PEN SUBCUTANEOUS	
TAFINLAR ORAL CAPSULE .....	76	SOLUTION AUTO-INJECTOR 200	
TAFINLAR ORAL TABLET SOLUBLE	77	MG/2ML.....	148, 149
TAGRISSE .....	239	TREMFYA SUBCUTANEOUS	
TALVEY.....	320	SOLUTION PREFILLED SYRINGE	148,
TALZENNA .....	319	149	
TASIGNA ORAL CAPSULE 150 MG, 200		tretinoin external cream .....	345
MG, 50 MG.....	217	trientine hcl oral capsule 250 mg.....	358
TAVNEOS .....	36	TRUE COMFORT ALCOHOL PREP	
TAZVERIK.....	322	PADS PAD 70 % .....	168, 171

TRUE COMFORT INSULIN SYRINGE  
30G X 1/2..... 168, 171  
TRUE COMFORT INSULIN SYRINGE  
30G X 5/16..... 168, 171  
TRUE COMFORT INSULIN SYRINGE  
31G X 5/16..... 168, 171  
TRUE COMFORT INSULIN SYRINGE  
32G X 5/16..... 168, 171  
TRUE COMFORT PRO ALCOHOL PREP  
PAD 70 % ..... 168, 171  
TRUE COMFORT PRO INSULIN SYR  
30G X 1/2..... 168, 171  
TRUE COMFORT PRO INSULIN SYR  
30G X 5/16..... 168, 171  
TRUE COMFORT PRO INSULIN SYR  
31G X 5/16..... 168, 171  
TRUE COMFORT PRO INSULIN SYR  
32G X 5/16..... 168, 171  
TRUEPLUS INSULIN SYRINGE 28G X  
1/2 ..... 168, 171  
TRUEPLUS INSULIN SYRINGE 29G X  
1/2 ..... 168, 171  
TRUEPLUS INSULIN SYRINGE 30G X  
5/16 ..... 168, 171  
TRUEPLUS INSULIN SYRINGE 31G X  
5/16 ..... 168, 171  
TRULICITY SUBCUTANEOUS  
SOLUTION AUTO-INJECTOR ..... 144  
TRUQAP ORAL TABLET ..... 63  
TRUQAP TABLET THERAPY PACK 160  
MG ORAL ..... 63  
TRUXIMA..... 287  
TUKYSA ORAL TABLET 150 MG, 50  
MG ..... 361  
TURALIO ..... 254  
TYENNE..... 339, 340, 341, 342  
U  
UBRELVY..... 362  
ULTICARE INSULIN SAFETY SYR 29G  
X 1/2..... 168, 171  
ULTICARE INSULIN SYRINGE 28G X  
1/2 ..... 168, 171  
ULTICARE INSULIN SYRINGE 29G X  
1/2 ..... 168, 169, 171  
ULTICARE INSULIN SYRINGE 30G X  
1/2 ..... 169, 171  
ULTICARE INSULIN SYRINGE 30G X  
5/16 ..... 169, 171  
ULTICARE INSULIN SYRINGE 31G X  
1/4 ..... 169, 171  
ULTICARE INSULIN SYRINGE 31G X  
5/16 ..... 169, 171  
ULTIGUARD SAFEPACK SYR/NEEDLE  
30G X 1/2..... 169, 171  
ULTIGUARD SAFEPACK SYR/NEEDLE  
31G X 5/16..... 169, 171  
ULTILET ALCOHOL SWABS PAD ... 169,  
171  
ULTILET INSULIN SYRINGE 30G X 1/2  
..... 169, 171  
ULTILET INSULIN SYRINGE 30G X 5/16  
..... 169, 171  
ULTILET INSULIN SYRINGE 31G X 1/4  
..... 169, 171  
ULTILET INSULIN SYRINGE 31G X  
15/64 ..... 169, 171  
ULTILET INSULIN SYRINGE 31G X 5/16  
..... 169, 171  
ULTILET INSULIN SYRINGE SHORT  
30G X 1/2..... 169, 171  
ULTILET INSULIN SYRINGE SHORT  
30G X 5/16..... 169, 171  
ULTILET INSULIN SYRINGE SHORT  
31G X 5/16..... 169, 171  
ULTRA COMFORT INSULIN SYRINGE  
30G X 5/16..... 169, 171  
ULTRA FLO INSULIN SYR 1/2 UNIT  
30G X 1/2..... 170, 171  
ULTRA FLO INSULIN SYR 1/2 UNIT  
30G X 5/16..... 170, 171  
ULTRA FLO INSULIN SYR 1/2 UNIT  
31G X 5/16..... 170, 171  
ULTRA FLO INSULIN SYRINGE 29G X  
1/2 ..... 170, 171  
ULTRA FLO INSULIN SYRINGE 30G X  
1/2 ..... 170, 171  
ULTRA FLO INSULIN SYRINGE 30G X  
5/16 ..... 170, 171  
ULTRA FLO INSULIN SYRINGE 31G X  
5/16 ..... 170, 171  
ULTRACARE INSULIN SYRINGE 30G X  
1/2 ..... 170, 171

ULTRACARE INSULIN SYRINGE 30G X 5/16 .....	170, 171	VIVIMUSTA .....	46
ULTRACARE INSULIN SYRINGE 31G X 5/16 .....	170, 171	VIZIMPRO .....	78
ULTRA-COMFORT INSULIN SYRINGE 29G X 1/2.....	170, 171	VONJO .....	241
ULTRA-THIN II INS SYR SHORT 30G X 5/16 .....	170, 171	VORANIGO .....	386
ULTRA-THIN II INS SYR SHORT 31G X 5/16 .....	170, 171	voriconazole oral suspension reconstituted .....	387
ULTRA-THIN II INSULIN SYRINGE 29G X 1/2.....	170, 171	VOSEVI.....	307
UPTRAVI INTRAVENOUS.....	298	VOWST .....	126
UPTRAVI ORAL TABLET 1000 MCG, 1200 MCG, 1400 MCG, 1600 MCG, 200 MCG, 400 MCG, 600 MCG, 800 MCG .....	298	VP INSULIN SYRINGE 29G X 1/2 .....	171
UPTRAVI TITRATION.....	298	VUMERITY .....	92
<b>V</b>		VYALEV SUBCUTANEOUS SOLUTION 12-240 MG/ML.....	134
VALCHLOR.....	204	VYLOY.....	391
VALUE HEALTH INSULIN SYRINGE 29G X 1/2.....	170, 171	<b>W</b>	
VANFLYTA .....	264	WEBCOL ALCOHOL PREP LARGE PAD 70 % .....	171
VANISHPOINT INSULIN SYRINGE 29G X 5/16.....	170, 171	WELIREG.....	45
VANISHPOINT INSULIN SYRINGE 30G X 3/16.....	170, 171	WINREVAIR.....	313
VANISHPOINT INSULIN SYRINGE 30G X 5/16.....	170, 171	<b>X</b>	
VEGZELMA.....	50	XALKORI ORAL CAPSULE.....	74
VENCLEXTA ORAL TABLET 10 MG, 100 MG, 50 MG.....	381	XALKORI ORAL CAPSULE SPRINKLE 150 MG, 20 MG, 50 MG .....	75
VENCLEXTA STARTING PACK .....	381	XDEMVI .....	198
VEOZAH .....	130	XELJANZ.....	343, 344
VERIFINE INSULIN SYRINGE 29G X 1/2 .....	170, 171	XELJANZ XR .....	343, 344
VERIFINE INSULIN SYRINGE 31G X 5/16 .....	170, 171	XERMELO .....	325
VERQUVO .....	382	XGEVA.....	86
VERZENIO.....	5	XIFAXAN ORAL TABLET 200 MG, 550 MG .....	274
vigabatrin .....	383	XOLAIR .....	236, 238
vigadrone.....	383	XOSPATA .....	141
vigpoder .....	383	XPOVIO (100 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 50 MG .....	299
VITRAKVI ORAL CAPSULE 100 MG, 25 MG .....	182	XPOVIO (40 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 10 MG, 40 MG .....	299
VITRAKVI ORAL SOLUTION .....	182	XPOVIO (40 MG TWICE WEEKLY) ORAL TABLET THERAPY PACK 40 MG .....	299
		XPOVIO (60 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 60 MG..	299
		XPOVIO (60 MG TWICE WEEKLY)...	299
		XPOVIO (80 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 40 MG..	299
		XPOVIO (80 MG TWICE WEEKLY)...	299

XTANDI ORAL CAPSULE.....	111, 112
XTANDI ORAL TABLET 40 MG, 80 MG .....	111, 112
XYOSTED.....	329
<b>Y</b>	
YERVOY.....	176
YESINTEK.....	373, 374, 375, 376
YONSA.....	7
YUFLYMA (1 PEN).....	13, 14, 15
YUFLYMA (2 SYRINGE).....	13, 14, 15
YUFLYMA-CD/UC/HS STARTER .	13, 14, 15
<b>Z</b>	
ZEJULA ORAL CAPSULE .....	221

ZEJULA ORAL TABLET.....	221
ZELBORAF.....	380
ZEVRX STERILE ALCOHOL PREP PAD PAD 70 % .....	171
ZIIHERA.....	388
ZIRABEV .....	52
ZOLADEX.....	147
ZTALMY .....	139
ZURZUVAE ORAL CAPSULE 20 MG, 25 MG, 30 MG.....	392
ZYDELIG .....	152
ZYKADIA ORAL TABLET .....	66
ZYNLONTA.....	196
ZYNYZ.....	270