Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: adalimumab-aacf (CF) Crohn 40 mg | | INDICATION: Humira Biosimilar FDA approved to treat seven inflammatory diseases including moderate-to-severe rheumatoid arthritis in adults, moderate-to-severe polyarticular juvenile idiopathic arthritis in patients 2 years of age and older, psoriatic arthritis in adults, ankylosing spondylitis in adults, moderate-to-severe chronic plaque psoriasis in adults, moderate-to-severe Crohn's disease in adults and pediatric patients 6 years of age and older and moderate-to-severe ulcerative colitis in adults |
|---|---------------|--|
| REASON FOR CHANGE: New Drug | | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Non-Formulary | Prior Authorization (CED), Quantity Limit |
| STANDARD FORMULARY | Non-Formulary | Quantity Limit |
| EXCHANGE FORMULARY | Non-Formulary | Quantity Limit |
| FAMIS FORMULARY | Non-Formulary | Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization (PDL Criteria), Quantity Limit |
| MEDICARE FORMULARY | Non-Formulary | N/A |
| OHANTITY LIMIT: N/A | | |

QUANTITY LIMIT: N/A

- (COMMERCIAL): 3 kits per 365 days
- (MEDICAID): 3 kits per 365 days
- (MEDICARE): N/A

FORMULARY ALTERNATIVES: (COMMERCIAL) Humira pen/syringe (Abbvie mfg only), Cyltezo (adalimumab-adbm), Yuflyma (adalimumab-aaty); **HIX/SG 2024 & 25** – Simlandi (adalimumab-ryvk) and adalimumab-adbm; (MEDICAID) Humira pen/syringe (Abbvie mfg only); (MEDICARE) Humira pen/syringe (Abbvie mfg only), Cyltezo (adalimumab-adbm), Yuflyma (adalimumab-aaty)

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: adalimumab-aacf (CF) PS-UV 40 mg | | INDICATION: Humira Biosimilar FDA approved to treat seven inflammatory diseases including moderate-to-severe rheumatoid arthritis in adults, moderate-to-severe polyarticular juvenile idiopathic arthritis in patients 2 years of age and older, psoriatic arthritis in adults, ankylosing spondylitis in adults, moderate-to-severe chronic plaque psoriasis in adults, moderate-to-severe Crohn's disease in adults and pediatric patients 6 years of age and older and moderate-to-severe ulcerative colitis in adults |
|---|---------------|--|
| REASON FOR CHANGE: New I | Orug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Non-Formulary | Prior Authorization (CED), Quantity Limit |
| STANDARD FORMULARY | Non-Formulary | Quantity Limit |
| EXCHANGE FORMULARY | Non-Formulary | Quantity Limit |
| FAMIS FORMULARY | Non-Formulary | Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization (PDL Criteria), Quantity Limit |
| MEDICARE FORMULARY | Non-Formulary | N/A |

QUANTITY LIMIT:

- (COMMERCIAL) 2 kits per 365 days
- (MEDICAID): 2 kits per 365 days
- (MEDICARE): N/A

FORMULARY ALTERNATIVES: (COMMERCIAL) Humira pen/syringe (Abbvie mfg only), Cyltezo (adalimumab-adbm), Yuflyma (adalimumab-aaty); **HIX/SG 2024 & 25** – Simlandi (adalimumab-ryvk) and adalimumab-adbm; (MEDICAID) Humira pen/syringe (Abbvie mfg only); (MEDICARE) Humira pen/syringe (Abbvie mfg only), Cyltezo (adalimumab-adbm), Yuflyma (adalimumab-aaty)

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: Aqneursa [™] (levacetylleucine) for oral suspension: 1-gram levacetylleucine in a unit-dose packet | | INDICATION: Treatment of neurological manifestations of Niemann-Pick disease type C (NPC) in adults and pediatric patients weighing ≥15 kg |
|---|--------------------|--|
| REASON FOR CHANGE: New | Drug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization, Quantity Limit |
| MEDICARE FORMULARY Specialty (Tier 5) | | Prior Authorization, Quantity Limit |
| QUANTITY LIMIT: 4 grams per day | | |
| FORMULARY ALTERNATIVES: N/A | | |

| DRUG NAME: Augtyro™ (repotrectinib) 40 mg capsules | | INDICATION: For the treatment of adult patients with locally advanced or metastatic ROS1-positive non-small cell lung cancer (NSCLC); Adult and pediatric patients 12 years of age and older with solid tumors that have a neurotrophic tyrosine receptor kinase (NTRK) gene fusion and are locally advanced or metastatic or where surgical resection is likely to result in severe morbidity; or have progressed following treatment or have no satisfactory alternative therapy |
|--|--------------------|--|
| REASON FOR CHANGE: Change Drug Tier and Quantity Limit | | Intity Limit |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| SENTARA COMMUNITY PLAN Formulary (MEDICAID) FORMULARY | | Prior Authorization, Quantity Limit |
| MEDICARE FORMULARY Specialty (Tier 5) | | Prior Authorization, Quantity Limit |
| QUANTITY LIMIT: • (COMMERCIAL): 6 capsules per day | | |

• (MEDICARE): N/A
FORMULARY ALTERNATIVES: N/A

(MEDICAID): 6 capsules per day

Effective: April 1, 2025

| DRUG NAME: Augtyro™ (repotrectinib) 160 mg capsules REASON FOR CHANGE: New Drug | | INDICATION: For the treatment of adult patients with locally advanced or metastatic ROS1-positive non-small cell lung cancer (NSCLC); Adult and pediatric patients 12 years of age and older with solid tumors that have a neurotrophic tyrosine receptor kinase (NTRK) gene fusion and are locally advanced or metastatic or where surgical resection is likely to result in severe morbidity; or have progressed following treatment or have no satisfactory alternative therapy |
|--|--------------------|--|
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY Formulary | | Prior Authorization, Quantity Limit |
| MEDICARE FORMULARY Specialty (Tier 5) | | Prior Authorization, Quantity Limit |
| QUANTITY LIMIT: 2 capsules per day | | |
| FORMULARY ALTERNATIVES: N/A | | |

| DRUG NAME: Aurlumyn™ (iloprost) injection concentrate, for IV use 100 mcg/mL | | INDICATION: For the treatment of severe frostbite in adults to reduce the risk of digit amputation |
|---|-----------------|---|
| REASON FOR CHANGE: New D |)rug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Medical Benefit | N/A |
| STANDARD FORMULARY | Medical Benefit | N/A |
| EXCHANGE FORMULARY | Medical Benefit | N/A |
| FAMIS FORMULARY | Medical Benefit | N/A |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Medical Benefit | N/A |
| MEDICARE FORMULARY | Medical Benefit | N/A |
| QUANTITY LIMIT: N/A | | |
| FORMULARY ALTERNATIVES: N/A | | |

Effective: April 1, 2025

| DRUG NAME: carbamazepine 200 mg chewable tablets | | INDICATION: For use as monotherapy in the acute treatment of hypomania and mild to moderate mania or episodes with mixed features associated with bipolar disorder; For use as monotherapy and adjunctive therapy in the treatment of patients with focal onset seizures and generalized onset seizures; For the treatment of trigeminal or glossopharyngeal neuralgia |
|--|---------------|--|
| REASON FOR CHANGE: New I | orug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Non-Formulary | Prior Authorization (CED) |
| STANDARD FORMULARY | Non-Formulary | N/A |
| EXCHANGE FORMULARY | Non-Formulary | N/A |
| FAMIS FORMULARY | Non-Formulary | N/A |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Formulary | N/A |
| MEDICARE FORMULARY Non-Formulary | | N/A |
| QUANTITY LIMIT: N/A | | |
| FORMULARY ALTERNATIVES: (COMMERCIAL) carbamazepine 100 mg chewable tablets; (MEDICA carbamazepine 100 mg chewable tablets | | rbamazepine 100 mg chewable tablets; (MEDICARE) |

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: Cobenfy™ (xanomeline and trospium hydrochloride) capsules (all strengths) | | INDICATION: For the treatment of schizophrenia in adults |
|---|--------------------|---|
| REASON FOR CHANGE: New D |)rug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Tier 3 | Step-Edit, Quantity Limit |
| STANDARD FORMULARY | Non-Formulary | Quantity Limit |
| EXCHANGE FORMULARY | Non-Formulary | Quantity Limit |
| FAMIS FORMULARY | Non-Formulary | Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization (PDL Criteria), Quantity Limit |
| MEDICARE FORMULARY | Specialty (Tier 5) | Prior Authorization, Quantity Limit |

QUANTITY LIMIT:

- (COMMERCIAL):
 - 2 capsules per day (all strengths)
 - 56 capsules (1 starter pack) per 365 days
- (MEDICAID):
 - 2 capsules per day (all strengths)
 - 56 capsules (1 starter pack) per 365 days
- (MEDICARE):
 - 2 capsules per day (all strengths)
 - 56 capsules (1 starter pack) per 180 days

FORMULARY ALTERNATIVES: (COMMERCIAL): aripiprazole tablets, clozapine tablets, olanzapine tablets, quetiapine IR/ER tablets, risperidone tablets/solution, ziprasidone capsules; (MEDICAID): aripiprazole tab, clozapine tab, lurasidone, olanzapine ODT/tab/IM, quetiapine fumarate ER, quetiapine tab, risperidone ODT/ soln/ tab, Vraylar™, ziprasidone cap

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: Crexont (carbidopa and levodopa) extended-release capsules, for oral use (all strengths) 35 mg / 140 mg, 52.5 mg / 210 mg, 70 mg / 280 mg, 87. 5 mg / 350 mg | | INDICATION: For the treatment of Parkinson's disease, post-encephalitic parkinsonism, and parkinsonism that may follow carbon monoxide intoxication or manganese intoxication in adults |
|---|---------------|--|
| REASON FOR CHANGE: New Drug | | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Non-Formulary | Prior Authorization (CED), Quantity Limit |
| STANDARD FORMULARY | Non-Formulary | Quantity Limit |
| EXCHANGE FORMULARY | Non-Formulary | Quantity Limit |
| FAMIS FORMULARY | Non-Formulary | Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Quantity Limit |
| MEDICARE FORMULARY | Non-Formulary | N/A |

QUANTITY LIMIT: N/A

- (COMMERCIAL): 6 capsules per day (all strengths)
- (MEDICAID): 6 capsules per day (all strengths)
- (MEDICARE): N/A

FORMULARY ALTERNATIVES: carbidopa-levodopa ER 25-100 & 50-200 mg tablets

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: Ebglyss™ (lebrikizumab-lbkz) 250 mg/2 mL single-dose prefilled pen/syringe with needle shield | | INDICATION: For the treatment of moderate to severe atopic dermatitis in adults and pediatric patients ≥12 years of age weighing ≥40 kg whose disease is not adequately controlled with topical prescription therapies or when those therapies are not advisable |
|--|--------------------|--|
| REASON FOR CHANGE: New Drug | | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization (PDL Criteria), Quantity Limit |
| MEDICARE FORMULARY | Non-Formulary | N/A |

QUANTITY LIMIT:

(COMMERCIAL): 1 injection per 28 days(MEDICAID): 1 injection per 28 days

(MEDICARE): N/A

FORMULARY ALTERNATIVES: (MEDICAID): Adbry[™], Dupixent[®], (both require prior authorization); (MEDICARE): Dupixent[®] (requires prior authorization)

| DRUG NAME: Femlyv [™] (ethinyl estradiol and norethindrone acetate ODT) 1 mg – 0.02 mg | | INDICATION: For pregnancy prevention |
|--|---------------|--------------------------------------|
| REASON FOR CHANGE: New Drug | | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Non-Formulary | Prior Authorization (CED) |
| STANDARD FORMULARY | Non-Formulary | N/A |
| EXCHANGE FORMULARY | Non-Formulary | N/A |
| FAMIS FORMULARY | Non-Formulary | N/A |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | N/A |
| MEDICARE FORMULARY | Non-Formulary | N/A |
| QUANTITY LIMIT: N/A | | |
| FORMULARY ALTERNATIVES: Aurovela, Junel, Larin, Loestrin, Microgestin, norethindrone-ethinyl estradiol tablets | | |

Effective: April 1, 2025

| DRUG NAME: FreeStyle Libre 2 plus sensors | | INDICATION: The latest innovation to the existing |
|--|-----------|--|
| | | FreeStyle Libre 2 system with the following new |
| | | features/updates from the FreeStyle Libre 2 sensor: |
| | | Extends the sensor wear up to 15 days, can work with insulin pumps & expands the age indication to |
| | | 2 years and older. The FreeStyle Libre 2 Plus |
| | | sensor is compatible with the current FreeStyle |
| | | Libre 2 app and FreeStyle Libre 2 reader |
| REASON FOR CHANGE: New | Drug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Tier 2 | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Tier 2 | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY | Tier 2 | Prior Authorization, Quantity Limit |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| MEDICARE FORMULARY Medical Part B Benefit | | Prior Authorization, Quantity Limit |
| QUANTITY LIMIT: 2 sensors per 30 days | | |
| FORMULARY ALTERNATIVES: N/A | | |

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: Hyrimoz (adalimumab-adaz) (CF) all strengths & formulations | | INDICATION: Humira Biosimilar FDA approved to treat nine inflammatory diseases including moderate-to-severe rheumatoid arthritis in adults, moderate-to-severe polyarticular juvenile idiopathic arthritis in patients 2 years of age and older, psoriatic arthritis in adults, ankylosing spondylitis in adults, moderate-to-severe chronic plaque psoriasis in adults, moderate-to-severe Crohn's disease in adults and pediatric patients 6 years of age and older, moderate-to-severe ulcerative colitis in adults, moderate-to-severe chronic plaque psoriasis in adults, moderate-to-severe hidradenitis suppurativa in adults, treatment of non-infectious intermediate, posterior, and panuveitis in adults |
|--|---------------|---|
| REASON FOR CHANGE: Chang | | T |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Non-Formulary | Prior Authorization (CED), Quantity Limit |
| STANDARD FORMULARY | Non-Formulary | Quantity Limit |
| EXCHANGE FORMULARY | Non-Formulary | Quantity Limit |
| FAMIS FORMULARY | Non-Formulary | Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | | Prior Authorization (PDL Criteria), Quantity Limit |
| MEDICARE FORMULARY | Non-Formulary | N/A |
| QUANTITY LIMIT: N/A | | |
| | | Humira pen/syringe (Abbvie mfg only), Cyltezo X/SG 2024 & 25 – Simlandi (adalimumab-ryvk) and |

adalimumab-adbm; (MEDICAID): Humira pen/syringe (Abbvie mfg only); (MEDICARE): Humira pen/syringe

(Abbvie mfg only), Cyltezo (adalimumab-adbm), Yuflyma (adalimumab-aaty)

Effective: April 1, 2025

| DRUG NAME: Itovebi™ (inavolisib) tablets, all strengths | | INDICATION: For use in combination with palbociclib and fulvestrant for the treatment of adults with endocrine-resistant, PIK3CA-mutated, hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative, locally advanced or metastatic breast cancer, as detected by an FDA-approved test, following recurrence on or aftercompleting adjuvant endocrine therapy |
|--|--------------------|--|
| REASON FOR CHANGE: New | Drug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| MEDICARE FORMULARY | Specialty (Tier 5) | Prior Authorization, Quantity Limit |
| QUANTITY LIMIT: • 3 mg – 2 tablets per day • 9 mg – 1 tablet per day | | |
| FORMULARY ALTERNATIVES: N/A | | |
| | | |

| DRUG NAME: Lagevrio (monupiravir) 200 mg capsules | | INDICATION: An investigational medicine used to treat adults with mild-to-moderate COVID-19 who are at risk for progression to severe COVID-19 including hospitalization or death, and for whom other COVID-19 treatment options approved or authorized by the FDA are not accessible or |
|--|------------------|--|
| | | clinically appropriate |
| REASON FOR CHANGE: Exclud | ie Drug | 1 |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Excluded Benefit | N/A |
| STANDARD FORMULARY | Excluded Benefit | N/A |
| EXCHANGE FORMULARY | Excluded Benefit | N/A |
| FAMIS FORMULARY | Excluded Benefit | N/A |
| SENTARA COMMUNITY PLAN Excluded Benefit (MEDICAID) FORMULARY | | N/A |
| MEDICARE FORMULARY Excluded Benefit | | N/A |
| QUANTITY LIMIT: N/A | | |
| FORMULARY ALTERNATIVES: Paxlovid™ (nirmatrelvir/ritonavir) tablets | | |

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: Livdelzi (seladelpar) 10 mg capsules | | INDICATION: For the treatment of primary biliary cholangitis in combination with ursodeoxycholic acid (UDCA) in adults who have had an inadequate response to UDCA, or as monotherapy in patients unable to tolerate UDCA |
|---|--------------------|---|
| REASON FOR CHANGE: New | Drug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization, Quantity Limit |
| MEDICARE FORMULARY | Specialty (Tier 5) | Prior Authorization, Quantity Limit |
| QUANTITY LIMIT: 1 capsule per day | | |
| FORMULARY ALTERNATIVES: N/A | | |

| DRUG NAME: Lumryz™ (sodium oxybate) for extended-release oral suspension 4.5-6-7.5 mg starter pack | | indication: For the treatment of cataplexy or excessive daytime sleepiness (EDS) in patients 7 years of age and older with narcolepsy |
|---|--------------------|---|
| REASON FOR CHANGE: New D |)rug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization, Quantity Limit |
| MEDICARE FORMULARY Non-Formulary | | N/A |
| QUANTITY LIMIT: | | |

- (COMMERCIAL): 28 packets (1 pack) per 365 days
- (MEDICAID): 28 packets (1 pack) per 365 days
- (MEDICARE): N/A

FORMULARY ALTERNATIVES: (MEDICARE) Xyrem & sodium oxybate solution (both require prior authorization)

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: Miplyffa™ (arimoclomol) capsules, all strengths | | INDICATION: For use in combination with miglustat for the treatment of neurological manifestations of Niemann-Pick disease type C (NPC) in adult and pediatric patients 2 years of age and older |
|---|--------------------|--|
| REASON FOR CHANGE: New D |)rug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization, Quantity Limit |
| MEDICARE FORMULARY Specialty (Tier 5) | | Prior Authorization, Quantity Limit |
| QUANTITY LIMIT: 3 capsules per day (all strengths) | | |
| FORMULARY ALTERNATIVES: N/A | | |
| | | |

| DRUG NAME: Neffy (epinephrine nasal spray) 2 mg/0.1 mL of epinephrine per spray | | INDICATION: For emergency treatment of type I allergic reactions, including anaphylaxis, in adult and pediatric patients who weigh 30 kg or greater | |
|---|-----------------------------|---|--|
| REASON FOR CHANGE: New [| REASON FOR CHANGE: New Drug | | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS | |
| OPEN FORMULARY | Non-Formulary | Prior Authorization (CED) | |
| STANDARD FORMULARY | Non-Formulary | N/A | |
| EXCHANGE FORMULARY | Non-Formulary | N/A | |
| FAMIS FORMULARY | Non-Formulary | N/A | |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization (PDL Criteria) | |
| MEDICARE FORMULARY | Non-Formulary | N/A | |
| QUANTITY LIMIT: N/A | | | |
| FORMULARY ALTERNATIVES: (COMMERCIAL) epinephrine 0.3 mg injection; (MEDICIAD) epinephrine | | | |

0.3 mg (authorized generic EpiPen®), Brand Epipen®; (MEDICARE) epinephrine 0.3 mg injection

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: Nemluvio™ (nemolizumab-ilto) for injection 30 mg | | INDICATION: For the treatment of prurigo nodularis in adults |
|--|--------------------|---|
| REASON FOR CHANGE: New Drug | | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization (PDL Criteria), Quantity Limit |
| MEDICARE FORMULARY | Specialty (Tier 5) | Prior Authorization, Quantity Limit |
| | | |

QUANTITY LIMIT:

- (COMMERCIAL): 1 injection (30 mg) per 28 days
- (MEDICAID): 1 injection (30 mg) per 28 days
- (MEDICARE): 2 injections (60 mg) per 28 days

FORMULARY ALTERNATIVES: (MEDICAID) Dupixent® (requires prior authorization)

| DRUG NAME: Ocrevus Zunovo™ (ocrelizumab and hyaluronidase-ocsq) solution in a single-dose vial containing 20 mg ocrelizumab and 23,000 units hyaluronidase per 23 mL (40 mg and 1,000 units per | | INDICATION: For the treatment of primary progressive multiple sclerosis (MS) in adults and relapsing forms of MS, including clinically isolated syndrome, relapsing remitting disease, and active | |
|---|---------------------------|---|--|
| mL) REASON FOR CHANGE: New Drug | | secondary progressive disease in adults | |
| FORMULARY TIER | | UTILIZATION MANAGEMENT REQUIREMENTS | |
| OPEN FORMULARY | Medical Benefit | Prior Authorization | |
| STANDARD FORMULARY | Medical Benefit | Prior Authorization | |
| EXCHANGE FORMULARY | Medical Benefit | Prior Authorization | |
| FAMIS FORMULARY | Medical Benefit | Prior Authorization | |
| SENTARA COMMUNITY PLAN | Non-Formulary [Pharmacy] | Prior Authorization (PDL Criteria) | |
| (MEDICAID) FORMULARY | Medical Benefit [Medical] | Prior Authorization | |
| MEDICARE FORMULARY | Non-Formulary | Prior Authorization | |
| QUANTITY LIMIT: N/A | | | |
| FORMULARY ALTERNATIVES: (MEDICAID): Kesimpta® (requires ST); | | | |

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: oxycodone HCL abuse deterrent tablets (5, 10 & 30 mg) | | INDICATION: For the management of acute or chronic moderate to severe pain when the use of an opioid analgesic is appropriate and for which alternative treatments are inadequate |
|---|-------------------|--|
| REASON FOR CHANGE: New | Drug | · |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Non-Formulary | Prior Authorization (CED) |
| STANDARD FORMULARY | Non-Formulary | N/A |
| EXCHANGE FORMULARY | Non-Formulary | N/A |
| FAMIS FORMULARY | Non-Formulary | N/A |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization (PDL Criteria) |
| MEDICARE FORMULARY | Non-Formulary | N/A |
| QUANTITY LIMIT: N/A | | |
| FORMULARY ALTERNATIVES: | oxycodone IR 10 m | g tablets (*requires prior authorization) |
| DRUG NAME: Roxybond (oxycodone HCL abuse deterrent tablets) 10 mg | | INDICATION: For treatment of pain severe enough to require an opioid analgesic and for which alternative treatments are inadequate. It is formulated with SentryBond™ abuse-deterrent technology which reduces potential abuse by intranasal and intravenous routes. |
| REASON FOR CHANGE: New I | Drug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Non-Formulary | Prior Authorization (CED) |
| STANDARD FORMULARY | Non-Formulary | N/A |
| EXCHANGE FORMULARY | Non-Formulary | N/A |
| FAMIS FORMULARY | Non-Formulary | N/A |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization (PDL Criteria) |

N/A

15

(MEDICAID) FORMULARY MEDICARE FORMULARY

QUANTITY LIMIT: N/A

Non-Formulary

FORMULARY ALTERNATIVES: oxycodone IR 10 mg tablets (*requires prior authorization)

Effective: April 1, 2025

| DRUG NAME: Tecelra® (afamitresgene autoleucel) suspension for IV infusion | | INDICATION: For the treatment of adults with unresectable or metastatic synovial sarcoma who have received prior chemotherapy, are HLA-A*02:01P, -A*02:02P, -A*02:03P, or -A*02:06P positive and whose tumor expresses the MAGE-A4 antigen as determined by FDA-approved or cleared companion diagnostic devices |
|---|-----------------|--|
| REASON FOR CHANGE: New | Drug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Medical Benefit | Prior Authorization |
| STANDARD FORMULARY | Medical Benefit | Prior Authorization |
| EXCHANGE FORMULARY | Medical Benefit | Prior Authorization |
| FAMIS FORMULARY | Medical Benefit | Prior Authorization |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Medical Benefit | Prior Authorization |
| MEDICARE FORMULARY Medical Benefit | | Prior Authorization |
| QUANTITY LIMIT: N/A | | |
| FORMULARY ALTERNATIVES: | | |

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

DRUG NAME: Tecentriq Hybreza™ (atezolizumab/hyaluronidase-tqjs) subcutaneous injection 1875 mg/30,000 units

INDICATION: For the treatment (as a single agent) of unresectable or metastatic alveolar soft part sarcoma in adults; Treatment (in combination with bevacizumab) of unresectable or metastatic hepatocellular carcinoma in adults who have not received prior systemic therapy: Treatment (in combination with cobimetinib and vemurafenib) of BRAF V600 mutation-positive (as determined by an approved test) unresectable or metastatic melanoma in adults; Adjuvant treatment (as a single agent) following resection and platinum-based chemotherapy in adults with stage II to IIIA nonsmall cell lung cancer (NSCLC) whose tumors have PD-L1 expression on ≥1% of tumor cells, as determined by an approved test; First-line treatment (as a single agent) of metastatic NSCLC in adults whose tumors have high PD-L1 expression (PD-L1 stained ≥50% of tumor cells [TC] or PD-L1 stained tumor-infiltrating immune cells [IC] covering ≥10% of the tumor area), as determined by an approved test, and with no EGFR or ALK genomic tumor aberrations; First-line treatment (in combination with bevacizumab, paclitaxel, and carboplatin) of metastatic nonsquamous NSCLC in adults with no EGFR or ALK genomic tumor aberrations; First-line treatment (in combination with paclitaxel [protein bound] and carboplatin) of metastatic nonsquamous NSCLC in adults with no EGFR or ALK genomic tumor aberrations; Treatment (as a single agent) of metastatic NSCLC in adults with disease progression during or following platinum-containing chemotherapy (patients with EGFR or ALK genomic aberrations should have disease progression on approved NSCLC therapy for EGFR or ALK genomic tumor mutations prior to receiving atezolizumab/hvaluronidase: First-line treatment (in combination with carboplatin and etoposide) of extensive-stage small cell lung cancer in adults

REASON FOR CHANGE: New Drug

| TEMOSTI OT STATE THE BIAS | | |
|---------------------------|-----------------|-------------------------------------|
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Medical Benefit | Prior Authorization |
| STANDARD FORMULARY | Medical Benefit | Prior Authorization |
| EXCHANGE FORMULARY | Medical Benefit | Prior Authorization |
| FAMIS FORMULARY | Medical Benefit | Prior Authorization |

Effective: April 1, 2025

| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Medical Benefit | Prior Authorization |
|--|-----------------|---------------------|
| MEDICARE FORMULARY | Medical Benefit | Prior Authorization |
| QUANTITY LIMIT: N/A | | |
| FORMULARY ALTERNATIVES: N/A | | |

| DRUG NAME: Tofidence™ (tocilizumab-bavi) vial for intravenous administration, all strengths | | INDICATION: Biogen's manufacturer's first FDA-approved biosimilar to IV Actemra. For use in adult patients with moderately to severely active rheumatoid arthritis (RA) who have had an inadequate response to one or more disease-modifying anti-rheumatic drugs (DMARDs); Patients 2 years of age and older with active polyarticular juvenile idiopathic arthritis (PJIA); Patients 2 years of age and older with active systemic juvenile idiopathic arthritis (SJIA) | |
|---|-----------------|---|--|
| REASON FOR CHANGE: New Drug | | | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS | |
| OPEN FORMULARY | Medical Benefit | Prior Authorization | |
| STANDARD FORMULARY | Medical Benefit | Prior Authorization | |
| EXCHANGE FORMULARY | Medical Benefit | Prior Authorization | |
| FAMIS FORMULARY | Medical Benefit | Prior Authorization | |
| SENTARA COMMUNITY PLAN [Pharmacy] | | Prior Authorization (PDL Criteria) | |
| (MEDICAID) FORMULARY Medical Benefit [Medical] | | Prior Authorization | |
| MEDICARE FORMULARY | Medical Benefit | Prior Authorization | |
| QUANTITY LIMIT: N/A | | | |
| FORMULARY ALTERNATIVES: (MEDICAID) Enbrel® pen/sureclick/syringe/vial, Humira® pen/syringe (Abbvie mfg only), infliximab (generic Remicade®) | | | |

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: Tremfya (guselkumab) 200 mg/20 mL (10 mg/mL) solution in a single-dose vial for intravenous infusion | | INDICATION: For the treatment of adult patients with moderately to severely active ulcerative colitis (UC) |
|--|---------------------------|---|
| REASON FOR CHANGE: New | Drug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Medical Benefit | Prior Authorization |
| STANDARD FORMULARY | Medical Benefit | Prior Authorization |
| EXCHANGE FORMULARY | Medical Benefit | Prior Authorization |
| FAMIS FORMULARY | Medical Benefit | Prior Authorization |
| SENTARA COMMUNITY PLAN | Non-Formulary [Pharmacy] | Prior Authorization (PDL Criteria) |
| (MEDICAID) FORMULARY | Medical Benefit [Medical] | Prior Authorization |
| MEDICARE FORMULARY | Medical Benefit | Prior Authorization |
| QUANTITY LIMIT: N/A | | |
| FORMULARY ALTERNATIVES: (MEDICAID) Enbrel® pen/sureclick/syringe/vial, Humira® pen/syringe, infliximab (generic Remicade®) | | |

| DRUG NAME: Tremfya (guselkumab) 200 mg/2 mL in a single-dose prefilled pen & syringe | | INDICATION: For the treatment of adult patients with moderately to severely active ulcerative colitis (UC) | |
|---|-----------------------------|---|--|
| REASON FOR CHANGE: New D | REASON FOR CHANGE: New Drug | | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS | |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit | |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit | |
| EXCHANGE FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit | |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit | |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization (PDL Criteria), Quantity Limit | |
| MEDICARE FORMULARY | Non-Formulary | N/A | |

QUANTITY LIMIT:

- (COMMERCIAL): 2 pens/syringes per 28 days
- (MEDICAID): 2 pens/syringes per 28 days
- (MEDICARE): N/A

FORMULARY ALTERNATIVES: (MEDICIAD) Enbrel® pen/sureclick/syringe/vial, Humira® pen/syringe, infliximab (generic Remicade®); (MEDICARE) Humira pen/syringe (Abbvie mfg only), Cyltezo (adalimumabadbm), Yuflyma (adalimumab-aaty), Stelara, Skyrizi

Effective: April 1, 2025

| DRUG NAME: Tyenne® (tocilizumab-aazg) vial for intravenous administration, all strengths REASON FOR CHANGE: New Drug | | INDICATION: Biosimilar to Actemra®. For treatment of adult patients with moderately to severely active rheumatoid arthritis who have had an inadequate response to one or more Disease-Modifying Anti-Rheumatic Drugs (DMARDs); Adult patients with giant cell arteritis; Patients 2 years of age and older with active polyarticular juvenile idiopathic arthritis; and Patients 2 years of age and older with active systemic juvenile idiopathic arthritis |
|---|---------------------------|---|
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Medical Benefit | Prior Authorization |
| STANDARD FORMULARY | Medical Benefit | Prior Authorization |
| EXCHANGE FORMULARY | Medical Benefit | Prior Authorization |
| FAMIS FORMULARY | Medical Benefit | Prior Authorization |
| SENTARA COMMUNITY PLAN [Pharmacy] | | Prior Authorization (PDL Criteria) |
| (MEDICAID) FORMULARY | Medical Benefit [Medical] | Prior Authorization |
| MEDICARE FORMULARY | Specialty (Tier 5) | Prior Authorization |
| QUANTITY LIMIT: N/A | | |
| FORMULARY ALTERNATIVES: (MEDICAID) Enbrel® pen/sureclick/syringe/vial, Humira® pen/syringe (Abbvie mfg only), infliximab (generic Remicade®) | | |

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: Tyenne® (tocilizumab-aazg) 162 mg/0.9 mL prefilled syringe/auto-injector for subcutaneous administration REASON FOR CHANGE: New Drug | | INDICATION: Biosimilar to Actemra®. For treatment of adult patients with moderately to severely active rheumatoid arthritis who have had an inadequate response to one or more Disease-Modifying Anti-Rheumatic Drugs (DMARDs); Adult patients with giant cell arteritis; Patients 2 years of age and older with active polyarticular juvenile idiopathic arthritis; and Patients 2 years of age and older with active systemic juvenile idiopathic arthritis |
|--|--------------------|---|
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization (PDL Criteria), Quantity Limit |
| MEDICARE FORMULARY | Non-Formulary | N/A |
| OLIANTITY LIMIT. | | |

QUANTITY LIMIT:

- (COMMERCIAL): 3.6 mL (4 injections) per 28 days
- (MEDICAID): 3.6 mL (4 injections) per 28 days
- (MEDICARE): N/A

FORMULARY ALTERNATIVES: (MEDICAID) Enbrel® pen/sureclick/syringe/vial, Humira® pen/syringe (Abbvie mfg only), infliximab (generic Remicade®); (MEDICARE) Actemra (*requires prior authorization)

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: Tryvio™ (aprocitentan) 12.5 mg tablets | | INDICATION: For use in patients whose BP is not adequately controlled on other antihypertensive medications |
|--|--------------------|--|
| REASON FOR CHANGE: New D | Drug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization, Quantity Limit |
| MEDICARE FORMULARY | Non-Formulary | N/A |

QUANTITY LIMIT:

• (COMMERCIAL): 1 tablet per day

• (MEDICAID): 1 tablet per day

• (MEDICARE): N/A

FORMULARY ALTERNATIVES: (MEDICARE) amlodipine tablets, valsartan tablets, hydrochlorothiazide tablets, spironolactone tablets

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: Undecatrex (testosterone undecanoate) 200 mg capsule | | INDICATION: For the treatment of testicular failure due to cryptorchidism, bilateral torsion, orchitis, vanishing testis syndrome, orchiectomy, Klinefelter syndrome, chemotherapy, or toxic damage from alcohol or heavy metals; gonadotropin or luteinizing hormone-releasing hormone deficiency; or pituitary-hypothalamic injury from tumors, trauma, or radiation |
|---|---------------|--|
| REASON FOR CHANGE: New D |)rug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Non-Formulary | Prior Authorization (CED) , Quantity Limit |
| STANDARD FORMULARY | Non-Formulary | Quantity Limit |
| EXCHANGE FORMULARY | Non-Formulary | Quantity Limit |
| FAMIS FORMULARY | Non-Formulary | Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Quantity Limit |
| MEDICARE FORMULARY | Non-Formulary | N/A |
| QUANTITY I IMIT: N/A | | |

QUANTITY LIMIT: N/A

• (COMMERCIAL): 2 capsules per day

• (MEDICAID): 2 capsules per day

• (MEDICARE): N/A

FORMULARY ALTERNATIVES: (COMMERCIAL) Kyzatrex 200 mg capsules, testosterone cypionate injection (*both require prior authorization); (MEDICIAD) testosterone cypionate injection (*requires prior authorization); (MEDICARE) testosterone cypionate injection (*requires prior authorization)

| DRUG NAME: Vyalev™ (foscarbiodpa and foslevodopa) injection for subcutaneous, injection contains 120 mg foscarbidopa and 2,400 mg foslevodopa per 10 mL (12 mg foscarbidopa and 240 mg foslevodopa per mL) | | INDICATION: For the treatment of motor fluctuations in adults with advanced Parkinson's disease |
|--|--------------------|---|
| REASON FOR CHANGE: New | Drug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization, Quantity Limit |
| MEDICARE FORMULARY | Specialty (Tier 5) | Prior Authorization, Quantity Limit |
| QUANTITY LIMIT: 6 cartons per 30 days | | |
| FORMULARY ALTERNATIVES: N/A | | |

Effective: April 1, 2025

| DRUG NAME: Vyloy (zolbetuximab-clzb) 100 mg lyophilized powder in a single-dose vial for IV infusion | | INDICATION: For use in combination with fluoropyrimidine- and platinum-containing chemotherapy for the first-line treatment of adults with locally advanced unresectable or metastatic human epidermal growth factor receptor 2 (HER2)-negative gastric or gastroesophageal junction adenocarcinoma whose tumors are claudin (CLDN) 18.2 positive as determined by an FDA-approved test |
|--|--------------------|---|
| REASON FOR CHANGE: New | Drug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Medical Benefit | Prior Authorization |
| STANDARD FORMULARY | Medical Benefit | Prior Authorization |
| EXCHANGE FORMULARY | Medical Benefit | Prior Authorization |
| FAMIS FORMULARY | Medical Benefit | Prior Authorization |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Medical Benefit | Prior Authorization |
| MEDICARE FORMULARY | Medical Benefit | Prior Authorization |
| QUANTITY LIMIT: N/A | | |
| FORMULARY ALTERNATIVES: N/A | | |
| DRUG NAME: Yorvipath (palopegteriparatide) single-patient use prefilled pen injection, for subcutaneous use (all strengths) | | INDICATION: For the treatment of hypoparathyroidism in adults |
| REASON FOR CHANGE: New | Drug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| FAMIS FORMULARY | Formulary | Prior Authorization, Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY Non-Formulary | | Prior Authorization, Quantity Limit |
| MEDICARE FORMULARY | Specialty (Tier 5) | Prior Authorization, Quantity Limit |
| QUANTITY LIMIT: 2 pens per 28 days (all strengths) | | |
| FORMULARY ALTERNATIVES: N/A | | |

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: Yuflyma® (adalimumab-aaty) (CF) all strengths & formulations | | INDICATION: Humira Biosimilar FDA approved to treat nine inflammatory diseases including moderate-to-severe rheumatoid arthritis in adults, moderate-to-severe polyarticular juvenile idiopathic arthritis in patients 2 years of age and older, psoriatic arthritis in adults, ankylosing spondylitis in adults, moderate-to-severe chronic plaque psoriasis in adults, moderate-to-severe Crohn's disease in |
|---|--------------------|--|
| REASON FOR CHANGE: Change Drug Tier and Utiliz | | adults and pediatric patients 6 years of age and older, moderate-to-severe ulcerative colitis in adults, moderate-to-severe chronic plaque psoriasis in adults, moderate-to-severe hidradenitis suppurativa in adults, treatment of non-infectious intermediate, posterior, and panuveitis in adults |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| STANDARD FORMULARY | Specialty (Tier 4) | Prior Authorization, Quantity Limit |
| EXCHANGE FORMULARY Non-Formulary | | Quantity Limit |

Prior Authorization, Quantity Limit

Prior Authorization, Quantity Limit

Prior Authorization (PDL Criteria), Quantity Limit

QUANTITY LIMIT:

FAMIS FORMULARY

SENTARA COMMUNITY PLAN

(MEDICAID) FORMULARY MEDICARE FORMULARY

- COMMERCIAL): 2 injections per 28 days
- (MEDICAID): 2 injections per 28 days
- (MEDICARE):
 - 20 & 80 mg: 2 injections per 28 days
 - 40 mg: 4 injections per 28 days

FORMULARY ALTERNATIVES: (MEDICAID): Humira pen/syringe (Abbvie mfg only)

Formulary

Non-Formulary

Specialty (Tier 5)

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

| DRUG NAME: Zituvimet (sitagliptin-metformin) IR tablets, all strengths | | INDICATION: For use as adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus |
|---|---------------|---|
| REASON FOR CHANGE: New | Drug | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS |
| OPEN FORMULARY | Non-Formulary | Prior Authorization (CED), Quantity Limit |
| STANDARD FORMULARY | Non-Formulary | Quantity Limit |
| EXCHANGE FORMULARY | Non-Formulary | Quantity Limit |
| FAMIS FORMULARY | Non-Formulary | Quantity Limit |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization (PDL Criteria), Quantity Limit |
| MEDICARE FORMULARY | Non-Formulary | N/A |
| QUANTITY LIMIT: N/A | | |

- COMMERCIAL): 2 tablets per day (both strengths)
- (MEDICAID): 2 tablets per day (both strengths)
- (MEDICARE): N/A

FORMULARY ALTERNATIVES: Janumet®

| DRUG NAME: Zituvimet XR (sitaç extended-release tablets, all stren | | INDICATION: For use as adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus | | | | | | |
|--|---------------|--|--|--|--|--|--|--|
| REASON FOR CHANGE: New D | | | | | | | | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS | | | | | | |
| OPEN FORMULARY | Non-Formulary | Prior Authorization (CED) | | | | | | |
| STANDARD FORMULARY | Non-Formulary | N/A | | | | | | |
| EXCHANGE FORMULARY | Non-Formulary | N/A | | | | | | |
| FAMIS FORMULARY | Non-Formulary | N/A | | | | | | |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization (PDL Criteria) | | | | | | |
| MEDICARE FORMULARY | Non-Formulary | N/A | | | | | | |
| QUANTITY LIMIT: N/A | | | | | | | | |
| FORMULARY ALTERNATIVES: Janumet® XR | | | | | | | | |

Effective: April 1, 2025

| DRUG NAME: Zituvio (sitagliptin) strengths | IR tablets, all | INDICATION: For use as adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus | | | | | | |
|---|-----------------|--|--|--|--|--|--|--|
| REASON FOR CHANGE: Change Drug Tier | | | | | | | | |
| FORMULARY | TIER | UTILIZATION MANAGEMENT REQUIREMENTS | | | | | | |
| OPEN FORMULARY | Non-Formulary | Prior Authorization (CED) | | | | | | |
| STANDARD FORMULARY | Non-Formulary | N/A | | | | | | |
| EXCHANGE FORMULARY | Non-Formulary | N/A | | | | | | |
| FAMIS FORMULARY | Non-Formulary | N/A | | | | | | |
| SENTARA COMMUNITY PLAN (MEDICAID) FORMULARY | Non-Formulary | Prior Authorization (PDL Criteria) | | | | | | |
| MEDICARE FORMULARY | Non-Formulary | N/A | | | | | | |
| QUANTITY LIMIT: N/A | | | | | | | | |
| FORMULARY ALTERNATIVES: Januvia® | | | | | | | | |

Effective: April 1, 2025

(For plans with pharmacy benefits administered by Sentara Health Plans)

4/1/2025 Commercial Formulary Updates

| Formulary Changes | | | | | | | | |
|--|-----------------------------------|----|---------------|--------------------------------|-----------------|---------------------------------|-------------------|--|
| APPLICABLE TO OTHER FORMULARIES (Y/N) | Label Name | SI | Drug Class | Current Formulary Status | Current Tier | Proposed Formulary Status | Propose d Tier | Preferred Alternatives for Clinically Equivalent Drugs (CED) & Non- Formulary (NF) Drugs |
| (CED/NF) ALL COMM EXCEPT SG2024 AND MDC CUSTOM | ACCUTANE 10 MG CAPSULE | Υ | F | Υ | 1 | N | 10 | AMNESTEEM 10 MG CAPSULE, CLARAVIS 10 MG CAPSULE, MYORISAN 10 MG CAPSULE, ZENATANE 10 MG CAPSULE |
| (CED/NF) ALL COMM EXCEPT SG2024 AND MDC CUSTOM | ACCUTANE 20 MG CAPSULE | Υ | F | Y | 1 | N | 10 | AMNESTEEM 20 MG CAPSULE, CLARAVIS 20 MG CAPSULE, MYORISAN 20 MG CAPSULE, ZENATANE 20 MG CAPSULE |
| (CED/NF) ALL COMM EXCEPT SG2024 AND MDC CUSTOM | ACCUTANE 30 MG CAPSULE | Υ | F | Υ | 1 | N | 10 | AMNESTEEM 30 MG CAPSULE, CLARAVIS 30 MG CAPSULE, MYORISAN 30 MG CAPSULE, ZENATANE 30 MG CAPSULE |
| (CED/NF) ALL COMM EXCEPT SG2024 AND MDC CUSTOM | ACCUTANE 40 MG CAPSULE | Υ | F | Υ | 1 | N | 10 | AMNESTEEM 40 MG CAPSULE, CLARAVIS 40 MG CAPSULE, MYORISAN 40 MG CAPSULE, ZENATANE 40 MG CAPSULE |
| (CED/NF) ALL COMM EXCEPT SG2024 AND MDC CUSTOM | DILTIAZEM 24H ER(LA) 420 MG TB | Υ | F | Υ | 2 | N | 10 | TIADYLT ER 420 MG CAPSULE, TIAZAC ER 420 MG CAPSULE |
| (CED/NF) ALL COMM EXCEPT SG2024 AND MDC CUSTOM | LIDOPIN 3% CREAM | Υ | F | Υ | 1 | N | 10 | LIDOCAINE 5% OINTMENT |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | AMABELZ 0.5 MG- 0.1 MG TABLET | Υ | F | Υ | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | AMABELZ 1 MG-0.5 MG TABLET | Υ | F | Υ | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | AMNESTEEM 10 MG CAPSULE | Υ | F | Υ | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | AMNESTEEM 20 MG CAPSULE | Υ | F | Υ | 1 | Y | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | AMNESTEEM 40 MG CAPSULE | Υ | F | Υ | 1 | Υ | 2 | N/A |

Effective: April 1, 2025

| APPLICABLE TO OTHER FORMULARIES (Y/N) | Label Name | SI | Drug Class | Current Formulary Status | Current Tier | Proposed Formulary Status | Propose d Tier | Preferred Alternatives for Clinically Equivalent Drugs (CED) & Non- Formulary (NF) Drugs |
|--|----------------------------------|----|---------------|--------------------------------|-----------------|---------------------------------|-------------------|--|
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | CLARAVIS 10 MG CAPSULE | Υ | F | Y | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | CLARAVIS 20 MG CAPSULE | Υ | F | Y | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | CLARAVIS 30 MG CAPSULE | Υ | F | Υ | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | CLARAVIS 40 MG CAPSULE | Υ | F | Υ | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | DOTTI 0.025 MG PATCH | Υ | F | Y | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | DOTTI 0.0375 MG PATCH | Υ | F | Y | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | DOTTI 0.05 MG PATCH | Υ | F | Υ | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | DOTTI 0.075 MG PATCH | Υ | F | Υ | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | DOTTI 0.1 MG PATCH | Υ | F | Υ | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | ESTRADIOL-NORETH 1-0.5 MG TAB | Υ | F | Υ | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | LYLLANA 0.025 MG PATCH | Υ | F | Υ | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | LYLLANA 0.0375 MG PATCH | Υ | F | Y | 1 | Υ | 2 | N/A |

Effective: April 1, 2025

| APPLICABLE TO OTHER FORMULARIES (Y/N) | Label Name | SI | Drug Class | Current Formulary Status | Current Tier | Proposed Formulary Status | Propose d Tier | Preferred Alternatives for Clinically Equivalent Drugs (CED) & Non- Formulary (NF) Drugs |
|--|---------------------------|----|---------------|--------------------------------|-----------------|---------------------------------|-------------------|--|
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | LYLLANA 0.05 MG PATCH | Υ | F | Υ | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | LYLLANA 0.075 MG PATCH | Υ | F | Y | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | LYLLANA 0.1 MG PATCH | Υ | F | Y | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | MYORISAN 10 MG CAPSULE | Υ | F | Y | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | MYORISAN 20 MG CAPSULE | Υ | F | Y | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | MYORISAN 30 MG CAPSULE | Y | F | Y | 1 | Y | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | MYORISAN 40 MG CAPSULE | Υ | F | Y | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | ZENATANE 10 MG CAPSULE | Y | F | Y | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | ZENATANE 20 MG CAPSULE | Υ | F | Υ | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | ZENATANE 30 MG CAPSULE | Υ | F | Υ | 1 | Υ | 2 | N/A |
| (T2) ALL COMM EXCEPT SG2024, VCUHS, COS, AND MDC CUSTOM | ZENATANE 40 MG CAPSULE | Υ | F | Υ | 1 | Υ | 2 | N/A |
| T4 ALL COMM EXCEPT MDC CUSTOM | EULEXIN 125 MG CAPSULE | X | F | N | 11 | Υ | 4 | N/A |

Effective: April 1, 2025

| APPLICABLE TO OTHER FORMULARIES (Y/N) | Label Name | SI | Drug Class | Current Formulary Status | Current Tier | Proposed Formulary Status | Propose d Tier | Preferred Alternatives for Clinically Equivalent Drugs (CED) & Non- Formulary (NF) Drugs |
|---|---------------------------------|----|---------------|--------------------------------|-----------------|---------------------------------|-------------------|--|
| T4 ALL COMM EXCEPT MDC CUSTOM | FLUTAMIDE 125 MG CAPSULE | Υ | F | Υ | 1 | Υ | 4 | N/A |
| T4 ALL COMM EXCEPT MDC CUSTOM | TOREMIFENE CITRATE 60 MG TAB | Υ | F | Υ | 2 | Υ | 4 | N/A |
| T4 OPEN/ & VCU ONLY (+ CHANGE) | FARESTON 60 MG TABLET | Х | F | Υ | 3 | Υ | 4 | N/A |
| T4 ALL CLOSED FORMULARIES EXCEPT MDC CUSTOM (+ CHANGE) | XERMELO 250 MG TABLET | W | F | N | 12 | Υ | 4 | N/A |