



NEW PREFERRED DRUGS	
THERAPEUTIC CLASS	NO PA REQUIRED PREFERRED
Central Nervous System (CNS) Agents: Anti-Migraine Agents, Acute	Sumatriptan Nasal Spray
Central Nervous System (CNS) Agents: Anticonvulsants*	Briviact
Central Nervous System (CNS) Agents: Atypical Antipsychotics*	Abilify Asimtufii
Central Nervous System (CNS) Agents: Multiple Sclerosis*	Kesimpta Teriflunomide
Endocrine Agents: Diabetes – Non-Insulin	Xigduo XR
Infectious Disease Agents: Antibiotics – Quinolones	Moxifloxacin
Infectious Disease Agents: Antivirals – HIV*	Apretude
Topical Agents: Antiparasitics	Vanalice

NEW CLINICAL PA REQUIRED PREFERRED DRUGS	
THERAPEUTIC CLASS	CLINICAL CRITERIA REQUIRED PREFERRED
Blood Formation, Coagulation, and Thrombosis Agents: Hemophilia Factor*	Jivi
Endocrine Agents: Androgens	Depo-Testosterone Testosterone Cypionate Testosterone Gel 1.62% Pump
Endocrine Agents: Growth Hormone	Zomacton
Respiratory Agents: Cystic Fibrosis	Trikafta Pak

NEW STEP THERAPY PREFERRED DRUGS	
THERAPEUTIC CLASS	STEP THERAPY REQUIRED PREFERRED
Central Nervous System (CNS) Agents: Atypical Antipsychotics*	Vraylar
Central Nervous System (CNS) Agents: Neuropathic Pain	Ztlido
Endocrine Agents: Diabetes – Insulin	Insulin Degludec
Ophthalmic Agents: Dry Eye Treatments	Xiidra

NEW NON PREFERRED DRUGS	
THERAPEUTIC CLASS	PA REQUIRED NON PREFERRED
Cardiovascular Agents: Angina, Hypertension and Heart Failure	Inpefa
Cardiovascular Agents: Pulmonary Arterial Hypertension*	Liqrev



Central Nervous System (CNS) Agents: Anti-Migraine Agents, Acute	Zavzpret
Central Nervous System (CNS) Agents: Atypical Antipsychotics*	Uzedy
Central Nervous System (CNS) Agents: Multiple Sclerosis*	Aubagio
Endocrine Agents: Androgens	Testosterone Gel 1% Pump
Endocrine Agents: Growth Hormone	Sogroya
Gastrointestinal Agents: Bowel Preparations	Suflave
Immunomodulator Agents: Systemic Inflammatory Disease	Adalimumab-adaz (Generic of Hyrimoz) Adalimumab-fkjp (Generic of Hudio) Cyltezo Hadlima Idacio Litfulo Yuflyma Yusimry
Infectious Disease Agents: Antivirals – HIV*	Cimduo Symfi Symfi Lo
Ophthalmic Agents: Dry Eye Treatments	Miebo
Respiratory Agents: Inhaled Agents	Tiotropium Inhaled Caps

THERAPEUTIC CATEGORIES WITH CHANGES IN CRITERIA	
Cardiovascular Agents: Angina, Hypertension & Heart Failure	
Central Nervous System (CNS) Agents: Attention Deficit Hyperactivity Disorder Agents	
Central Nervous System (CNS) Agents: Narcolepsy	
Central Nervous System (CNS) Agents: Neuropathic Pain	
Dermatologic Agents: Oral Acne Products	
Endocrine Agents: Diabetes – Hypoglycemia Treatments	
Endocrine Agents: Diabetes – Non-Insulin	
Endocrine Agents: Endometriosis	
Endocrine Agents: Uterine Fibroids	
Gastrointestinal Agents: Anti-Emetics	
Genitourinary Agents: Benign Prostatic Hyperplasia	
Immunomodulator Agents: Systemic Inflammatory Disease	
Infectious Disease Agents: Antifungals	
Respiratory Agents: Cystic Fibrosis	

REVISED THERAPEUTIC CATEGORY CRITERIA	
THERAPEUTIC CLASS	SUMMARY OF CHANGE



<p>Cardiovascular Agents: Angina, Hypertension & Heart Failure</p>	<p>ADDITIONAL SOTAGLIFLOZIN (INPEFA) CRITERIA:</p> <ul style="list-style-type: none"> Must provide documentation of an inadequate clinical response to at least <u>two</u> SGLT2 Inhibitors (refer to Endocrine Agents: Diabetes – Non-Insulin class for complete list)
<p>Central Nervous System (CNS) Agents: Attention Deficit Hyperactivity Disorder Agents</p>	<p>STEP THERAPY CRITERIA:</p> <ul style="list-style-type: none"> Must have had an inadequate clinical response of at least <u>30 days</u> with atomoxetine OR at least <u>two</u> preferred <u>stimulants-ADHD agents</u>. <p>AR – <u>Adderall</u>, <u>Amphetamine/Dextroamphetamine</u>, <u>Dexedrine-Dextroamphetamine IR</u>, & Zenedi: a PA is required for patients younger than 3 years</p> <p>AR – <u>Adderall XR</u>, <u>Amphetamine/Dextroamphetamine XR</u>, Atomoxetine, Cotempla XR-ODT, Daytrana, <u>Methylphenidate Patches</u>, <u>Dexedrine ER</u>, <u>Dextroamphetamine ER</u>, Dexamethylphenidate, Methylphenidate IR & ER, & Xelstrym: a PA is required for patients younger than 6 years</p>
<p>Central Nervous System (CNS) Agents: Narcolepsy</p>	<p>NON-PREFERRED CRITERIA:</p> <ul style="list-style-type: none"> Must have had an inadequate clinical response <u>of at least 30 days</u> with at least <u>two</u> preferred drugs - either (1) <u>at least 30 days</u> of modafinil or armodafinil; or (2) <u>at least 7 days of a</u> preferred methylphenidate or amphetamine drug
<p>Central Nervous System (CNS) Agents: Neuropathic Pain</p>	<p>STEP THERAPY CRITERIA:</p> <ul style="list-style-type: none"> Must have had an inadequate clinical response of at least <u>30 days</u> with generic Lidocaine patch
<p>Dermatologic Agents: Oral Acne Products</p>	<p>CLINICAL PA CRITERIA:</p> <ul style="list-style-type: none"> <u>Patient must be</u> Prescriber attests that patient is registered and meets all of the requirements of the iPLEDGE program
<p>Endocrine Agents: Diabetes – Hypoglycemia Treatments</p>	<p>NON-PREFERRED CRITERIA:</p> <ul style="list-style-type: none"> Must have had an inadequate clinical response of at least <u>two-one</u> preferred drugs OR the inability of the member and/or caregiver to administer a preferred glucagon product in a timely fashion
<p>Endocrine Agents: Diabetes – Non-Insulin</p>	<p>ADDITIONAL INFORMATION</p> <ul style="list-style-type: none"> An inadequate clinical response is defined as the inability to reach A1C goal after at least 120 days of current regimen, <u>with use of multiple-two or more drugs concomitantly per ADA guidelines, documented adherence, and appropriate dose escalation (must achieve maximum recommended dose or document that maximum recommended dose is not tolerated or is clinically inappropriate).</u> <ul style="list-style-type: none"> Must include a patient specific <u>document</u> A1C goal if less than <u>7%</u> <u>per ADA guidelines and</u> Must include current A1C (within last <u>6 months</u>)



	<ul style="list-style-type: none"> • Requests may be authorized for patients with a condition that is difficult to control (i.e., prone to ketoacidosis, hypoglycemia) <p>SUBSEQUENT AUTHORIZATION CRITERIA:</p> <ul style="list-style-type: none"> • Must provide documentation of patient’s clinical response to treatment and ongoing safety monitoring <ul style="list-style-type: none"> ○ Must document A1C goal per ADA guidelines and A1C trends including current value (within last 6 months). ○ Must include a patient specific A1C goal if less than 7% ○ Must include current A1C (within last 6 months) • Must meet all initial clinical criteria for subsequent authorizations.
<p>Endocrine Agents: Endometriosis</p>	<p>STEP THERAPY CRITERIA:</p> <ul style="list-style-type: none"> • Must have had an inadequate clinical response of at least 84 days with at least one preferred NSAID and one preferred oral contraceptive
<p>Endocrine Agents: Uterine Fibroids</p>	<p>CLINICAL PA CRITERIA:</p> <ul style="list-style-type: none"> • Must have had an inadequate clinical response of at least 90 days with at least one preferred oral contraceptive
<p>Gastrointestinal Agents: Anti-Emetics</p>	<p>NON-PREFERRED CRITERIA:</p> <ul style="list-style-type: none"> • Must have had an inadequate clinical response of at least 7-3 days with at least one preferred drug
<p>Genitourinary Agents: Benign Prostatic Hyperplasia</p>	<p>TADALAFIL (CIALIS) CRITERIA:</p> <ul style="list-style-type: none"> • Must have had an inadequate clinical response of at least 30 days with at least one alpha-1 adrenergic blocker. and If prostate volume of > 30cc on imaging, a prostate specific antigen (PSA) > 1.5ng/dL, or palpable prostate enlargement on digital rectal exam (DRE), then a trial of at least 90 days of finasteride is required.”
<p>Immunomodulator Agents: Systemic Inflammatory Disease</p>	<p>ALL AUTHORIZATIONS:</p> <ul style="list-style-type: none"> • First line treatment can vary based upon the severity of disease for certain diagnoses. Documentation of the patient’s disease state and the criteria used to classify the severity is required. <p>ADDITIONAL CROHN’S DISEASE CRITERIA:</p> <ul style="list-style-type: none"> • Must provide documentation of Crohn's Disease Activity Index (CDAI) score of 220 or higher to be considered moderate to severe disease <p>ADDITIONAL HIDRADENTIS SUPPURATIVA CRITERIA:</p> <ul style="list-style-type: none"> • Must provide documentation of Hurley Stage III to be classified as severe disease <p>ADDITIONAL PLAQUE PSORIASIS CRITERIA:</p> <ul style="list-style-type: none"> • To classify as severe disease patient must present at least two of



	the following: Psoriasis Area and Severity Index (PASI) score \geq 11, BSA \geq 10%, and Static Physician's Global Assessment (sPGA) \geq 3
Infectious Disease Agents: Antifungals	NON-PREFERRED CRITERIA: <ul style="list-style-type: none">• Must have had an inadequate clinical response of at least 7 days with at least <u>one-two</u> preferred drugs, if indicated for the diagnosis
Respiratory Agents: Cystic Fibrosis	AR – Trikafta Pak: a PA is required for patients 6 years and older

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