

NEW PREFERRED DRUGS	
THERAPEUTIC CLASS NO PA REQUIRED PREFERRED	
Cardiovascular Agents: Angina, Hypertension and Heart Failure	bisoprolol 5, 10mg labetalol 100, 200, 300mg spironolactone tab
Cardiovascular Agents: Antiarrhythmics	MULTAQ
Central Nervous System (CNS) Agents: Fibromyalgia Agents	SAVELLA
Central Nervous System (CNS) Agents: Neuropathic Pain	GRALISE HORIZANT
Central Nervous System (CNS) Agents: Skeletal Muscle Relaxants, Non-Benzodiazepine	methocarbamol 500, 750mg
Central Nervous System (CNS) Agents: Restless Legs Syndrome	HORIZANT
Endocrine Agents: Diabetes – Non-Insulin	glimepiride 1, 2, 4mg
Gastrointestinal Agents: Bowel Preparations	MOVIPREP
Gastrointestinal Agents: Crohn's Disease	mercaptopurine tab
Gastrointestinal Agents: Ulcerative Colitis	mesalamine ER cap 500mg PENTASA 250mg
Gastrointestinal Agents: Unspecified GI	polyethylene glycol oral powder bottle
Infectious Disease Agents: Antivirals – HIV* LEGACY CATEGORY	RUKOBIA VIREAD 150, 200mg
Respiratory Agents: Inhaled Agents	arformoterol neb
Topical Agents: Antifungals	tolnaftate cream, powder
Topical Agents: Immunomodulators	pimecrolimus [labeler 68682]

NEW CLINICAL PA REQUIRED PREFERRED DRUGS	
THERAPEUTIC CLASS CLINICAL CRITERIA REQUIRED PREFERRED	
Respiratory Agents: Cystic Fibrosis	ALYFTREK

NEW NON-PREFERRED DRUGS	
THERAPEUTIC CLASS	PA REQUIRED NON-PREFERRED



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Analgesic Agents: Opioids	tramadol IR 75mg
Blood Formation, Coagulation, and Thrombosis	HYMPAVZI
Agents: Hemophilia A, von Willebrand Disease,	
and Factor XIII Deficiency* LEGACY CATEGORY	
Blood Formation, Coagulation, and Thrombosis	HYMPAVZI
Agents: Hemophilia B* LEGACY CATEGORY	
Blood Formation, Coagulation, and Thrombosis	rivaroxaban tab
Agents: Oral Anticoagulants	
Blood Formation, Coagulation, and Thrombosis	ticagrelor
Agents: Oral Antiplatelet	1. 1.05
Cardiovascular Agents: Angina, Hypertension and Heart Failure	bisoprolol 2.5mg CORLANOR SOLN
Heart Failure	ivabradine tab
	labetalol 400mg
Cardiovascular Agents: Antiarrhythmics	quinidine IR, ER
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Central Nervous System (CNS) Agents: Alzheimer's Agents* LEGACY CATEGORY	memantine/donepezil cap 14-10, 21-10, 28-10mg
Central Nervous System (CNS) Agents:	VIGAFYDE
Anticonvulsants* LEGACY CATEGORY	VIGAL TOE
Central Nervous System (CNS) Agents: Atypical	EQUETRO
Antipsychotics* LEGACY CATEGORY	ERZOFRI
	OPIPZA
Central Nervous System (CNS) Agents:	gabapentin ER
Neuropathic Pain	
Central Nervous System (CNS) Agents:	VYALEV
Parkinson's Agents	
Central Nervous System (CNS) Agents: Skeletal	methocarbamol 1000mg
Muscle Relaxants, Non-Benzodiazepine	
Endocrine Agents: Androgens	AZMIRO
Endocrine Agents: Diabetes – Hypoglycemia	glucagon emerg kit labeler 00378
Treatments	
Endocrine Agents: Diabetes – Non-Insulin	glimepiride 3mg
	metformin IR 750mg
Gastrointestinal Agents: Ulcerative Colitis	ZITUVIMET XR
	PENTASA 500mg
Gastrointestinal Agents: Unspecified GI	polyethylene glycol oral powder packet
Conitarininami Apontos Electrichita Devilati	prucalopride
Genitourinary Agents: Electrolyte Depleter	ferric citrate tab
Agents Immunomodulator Agents: Systemic	NEMLUVIO
Inflammatory Disease	INLIVILUVIU
Infectious Disease Agents: Antibiotics –	cefaclor ER
Cephalosporins	CONTROL EN
Infectious Disease Agents: Antivirals – HIV*	EMTRIVA SOLN
LEGACY CATEGORY	VIREAD 250, 300mg TAB
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Ophthalmic Agents: Glaucoma Agents	BETIMOL 0.25%	
	timolol hemihydrate soln 0.5%	
Respiratory Agents: Epinephrine	epinephrine (labeler 00093, 00115)	
	NEFFY	
Respiratory Agents: Inhaled Agents	BROVANA	
	umeclidinium/vilanterol	
Topical Agents: Antifungals	tolnaftate soln	
Topical Agents: Immunomodulators	pimecrolimus [labeler 00591, 68462]	

THERAPEUTIC CATEGORIES WITH CHANGES IN CRITERIA
Analgesic Agents: Gout
Analgesic Agents: NSAIDS
Analgesic Agents: Opioids
Blood Formation, Coagulation, and Thrombosis Agents: Colony Stimulating Factors
Blood Formation, Coagulation, and Thrombosis Agents: Hematopoietic Agents
Blood Formation, Coagulation, and Thrombosis Agents: Hemophilia A, von Willebrand Disease, and
Factor XIII Deficiency* LEGACY CATEGORY
Blood Formation, Coagulation, and Thrombosis Agents: Hemophilia B* LEGACY CATEGORY
Blood Formation, Coagulation, and Thrombosis Agents: Heparin-Related Preparations
Blood Formation, Coagulation, and Thrombosis Agents: Oral Anticoagulants
Blood Formation, Coagulation, and Thrombosis Agents: Oral Antiplatelet
Cardiovascular Agents: Angina, Hypertension and Heart Failure
Cardiovascular Agents: Antiarrhythmics
Cardiovascular Agents: Lipotropics
Cardiovascular Agents: Pulmonary Arterial Hypertension* LEGACY CATEGORY
Central Nervous System (CNS) Agents: Alzheimer's Agents* LEGACY CATEGORY
Central Nervous System (CNS) Agents: Anticonvulsants* LEGACY CATEGORY
Central Nervous System (CNS) Agents: Anticonvulsants Rescue
Central Nervous System (CNS) Agents: Antidepressants* LEGACY CATEGORY
Central Nervous System (CNS) Agents: Attention Deficit Hyperactivity Disorder Agents
Central Nervous System (CNS) Agents: Atypical Antipsychotics* LEGACY CATEGORY
Central Nervous System (CNS) Agents: Fibromyalgia Agents
Central Nervous System (CNS) Agents: Medication Assisted Treatment of Opioid Addiction
Central Nervous System (CNS) Agents: Multiple Sclerosis* LEGACY CATEGORY
Central Nervous System (CNS) Agents: Narcolepsy
Central Nervous System (CNS) Agents: Neuropathic Pain
Central Nervous System (CNS) Agents: Parkinson's Agents
Central Nervous System (CNS) Agents: Restless Legs Syndrome
Central Nervous System (CNS) Agents: Sedative-Hypnotics, Non-Barbiturate
Central Nervous System (CNS) Agents: Skeletal Muscle Relaxants, Non-Benzodiazepine
Dermatologic Agents: Oral Acne Products
Dermatologic Agents: Topical Acne Products
Endocrine Agents: Androgens
Endocrine Agents: Diabetes – Hypoglycemia Treatments
Endocrine Agents: Diabetes – Insulin

Endocrine Agents: Diabetes - Non-Insulin

Endocrine Agents: Endometriosis Endocrine Agents: Estrogenic Agents Endocrine Agents: Growth Hormone

Endocrine Agents: Osteoporosis – Bone Ossification Enhancers

Gastrointestinal Agents: Anti-Emetics

Gastrointestinal Agents: Bowel Preparations
Gastrointestinal Agents: Crohn's Disease

Gastrointestinal Agents: Irritable Bowel Syndrome (IBS) with Diarrhea

Gastrointestinal Agents: Pancreatic Enzymes
Gastrointestinal Agents: Proton Pump Inhibitors

Gastrointestinal Agents: Ulcerative Colitis
Gastrointestinal Agents: Unspecified GI

Genitourinary Agents: Benign Prostatic Hyperplasia Genitourinary Agents: Electrolyte Depleter Agents

Genitourinary Agents: Urinary Antispasmodics

Hyperkalemia Agents: Potassium Binders

Immunomodulator Agents: Systemic Inflammatory Disease

Infectious Disease Agents: Antibiotics – Cephalosporins

Infectious Disease Agents: Antibiotics – Inhaled

Infectious Disease Agents: Antibiotics – Macrolides

Infectious Disease Agents: Antibiotics – Quinolones

Infectious Disease Agents: Antibiotics – Tetracyclines

Infectious Disease Agents: Antifungals

Infectious Disease Agents: Antivirals – Hepatitis C Agents

Infectious Disease Agents: Antivirals – Herpes

Infectious Disease Agents: Antivirals – HIV* LEGACY CATEGORY

Ophthalmic Agents: Antibiotic and Antibiotic-Steroid Combination Drops and Ointments

Ophthalmic Agents: Antihistamines & Mast Cell Stabilizers

Ophthalmic Agents: Dry Eye Treatments
Ophthalmic Agents: Glaucoma Agents

Ophthalmic Agents: NSAIDs

Ophthalmic Agents: Ophthalmic Steroids

Otic Agents: Antibacterial and Antibacterial/Steroid Combinations

Respiratory Agents: Antihistamines – Second Generation

Respiratory Agents: Cystic Fibrosis
Respiratory Agents: Epinephrine

Respiratory Agents: Hereditary Angioedema

Respiratory Agents: Inhaled Agents

Respiratory Agents: Leukotriene Receptor Modifiers & Inhibitors

Respiratory Agents: Monoclonal Antibodies-Anti-IL/Anti-IgE

Respiratory Agents: Nasal Preparations Respiratory Agents: Pulmonary Fibrosis

Topical Agents: Antifungals

Topical Agents: Antiparasitics

Topical Agents: Corticosteroids

Topical Agents: Immunomodulators



			
REVISED THERAPEUTIC CATEGORY CRITERIA			
THERAPEUTIC CLASS	SUMMARY OF CHANGE		
Analgesic Agents: Gout	 NON-PREFERRED CRITERIA: Must have had an inadequate clinical response of at least 30 days with at least one preferred drug in this UPDL category and indicated for diagnosis 		
Analgesic Agents: NSAIDS	 NON-PREFERRED CRITERIA: Must have had an inadequate clinical response of at least 30 da with at least two preferred drugs in this UPDL category, if and indicated for diagnosis 		
Analgesic Agents: Opioids	 NON-PREFERRED CRITERIA: Must have had an inadequate clinical response of at least 7 days of at least two unrelated preferred drugs with different active ingredients of the same duration of action (SHORT-ACTING or LONG-ACTING) AR − All codeine and tramadol containing products: a PA is required for patients younger than 12 years old 		
Blood Formation, Coagulation, and Thrombosis Agents: Colony Stimulating Factors Blood Formation, Coagulation, and	NON-PREFERRED CRITERIA: Must have had an inadequate clinical response of at least 14 days with at least one preferred drug in this UPDL category and indicated for diagnosis NON-PREFERRED CRITERIA: Must have had an inadequate clinical response of at least 14 days		
Thrombosis Agents: Hematopoietic Agents	with at least <u>one preferred</u> drug in this UPDL category <mark>and indicated for diagnosis</mark>		
Blood Formation, Coagulation, and Thrombosis Agents: Hemophilia A, von Willebrand Disease, and Factor XIII Deficiency* LEGACY CATEGORY	 NON-PREFERRED CRITERIA: Must have had an inadequate clinical response of at least 14 days with at least one preferred drug in this UPDL category and indicated for diagnosis ADDITIONAL HYMPAVZI (MARSTACIMAB-HNCQ) CRITERIA Must have had an inadequate clinical response of at least 30 days with HEMLIBRA 		
Blood Formation, Coagulation, and Thrombosis Agents: Hemophilia B* LEGACY CATEGORY	 NON-PREFERRED CRITERIA: Must have had an inadequate clinical response of at least 14 days with at least one preferred drug in this UPDL category and indicated for diagnosis 		
Blood Formation, Coagulation, and Thrombosis Agents: Heparin-Related Preparations	 NON-PREFERRED CRITERIA: Must have had an inadequate clinical response of at least 14 days with at least one preferred drug in this UPDL category and indicated for diagnosis 		



Blood Formation,	NON-PREFERRED CRITERIA:
Coagulation, and	 Must have had an inadequate clinical response of at least 14 days
Thrombosis Agents:	with at least two preferred drugs in this UPDL category and
Oral Anticoagulants	indicated for diagnosis
Blood Formation,	All products are covered without a PA
Coagulation, and	LENGTH OF AUTHORIZATION: 365 days
Thrombosis Agents:	LENGTH OF AUTHORIZATION. 503 days
_	NON PREFERRED CRITERIA
Oral Antiplatelet	NON-PREFERRED CRITERIA:
	 Must have had an inadequate clinical response of at least <u>30 days</u>
	with at least <u>one preferred</u> drug in this UPDL category and indicated
	for diagnosis
Cardiovascular	NON-PREFERRED CRITERIA:
Agents: Angina,	 Must have had an inadequate clinical response of at least <u>30 days</u>
Hypertension and	of at least two preferred drugs within the same sub-section
Heart Failure	classification in this UPDL category and indicated for diagnosis with
	the same mechanism of action, if available and indicated for the
	same diagnosis in this UPDL category
Cardiovascular	NON-PREFERRED CRITERIA:
Agents:	Must have had an inadequate clinical response of at least 30 days
Antiarrhythmics	with at least one preferred drug in this UPDL category and indicated
Antiarriytimics	
Cardiovascular	for diagnosis
	NON-PREFERRED CRITERIA:
Agents: Lipotropics	Must have had an inadequate clinical response of at least 30 days
	(or <u>90 days</u> for fibrates) with at least <u>one preferred</u> drug within the
	same sub-section classification in this UPDL category and indicated
	for diagnosis in the same drug class
Cardiovascular	NON-PREFERRED CRITERIA:
Agents: Pulmonary	 Must have had an inadequate clinical response of at least 30 days
Arterial	with at least <u>two preferred</u> drugs in this UPDL category <mark>and</mark>
Hypertension*	indicated for diagnosis, if available, one of which must be a
LEGACY CATEGORY	phosphodiesterase-5 inhibitor
Central Nervous	NON-PREFERRED CRITERIA:
System (CNS)	Must have had an inadequate clinical response of at least 30 days
Agents: Alzheimer's	with at least <u>two preferred</u> drugs in this UPDL category and
Agents* LEGACY	indicated for diagnosis
CATEGORY	
Central Nervous	NON-PREFERRED CRITERIA:
System (CNS)	Must have had an inadequate clinical response of at least 14 days
1	
Agents: Anti-	with at least <u>one</u> preferred drug and <u>one</u> step therapy drug in this
Migraine Agents,	UPDL category and indicated for diagnosis, if available one of which
Acute	has the same mechanism of action if available
Central Nervous	NON-PREFERRED CRITERIA:
System (CNS)	Must have had an inadequate clinical response of at least <u>60 days</u>
Agents: Anti-	to at least <u>one</u> <u>preferred</u> drug in this UPDL category <mark>and indicated</mark>
Migraine Agents,	for diagnosis
Cluster Headache	
Central Nervous	NON-PREFERRED CRITERIA:
System (CNS)	
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Agents: Anticonvulsants* LEGACY CATEGORY	 Must have had an inadequate clinical response of at least 30 days with at least two preferred drugs in this UPDL category and indicated for diagnosis For prescribers who are credentialed as a neurology specialty with Ohio Medicaid, there must have been an inadequate clinical response of at least 30 days with one preferred anticonvulsant drug in the standard tablet/capsule dosage form. Prescriptions submitted from a prescriber who is credentialed as a neurology specialty with Ohio Medicaid AND for drugs that are used only for seizures, there must have been an inadequate clinical response of at least 30 days with one preferred drug. This provision applies only to the standard tablet/capsule dosage form.
Central Nervous System (CNS) Agents: Anticonvulsants Rescue	AR – VALTOCO: a PA is required for patients younger than <mark>6</mark> 2 years old
Central Nervous System (CNS) Agents: Antidepressants* LEGACY CATEGORY	 NON-PREFERRED CRITERIA: Must have had an inadequate clinical response of at least 30 days with at least two preferred drugs in this UPDL category and indicated for diagnosis
Central Nervous System (CNS) Agents: Attention Deficit Hyperactivity Disorder Agents	 Must have had an inadequate clinical response of at least 30 days with at least two preferred drugs in this UPDL category and indicated for diagnosis.; if available
Central Nervous System (CNS) Agents: Atypical Antipsychotics* LEGACY CATEGORY	 NON-PREFERRED CRITERIA: Must have had an inadequate clinical response of at least 30 days with at least two preferred drugs in this UPDL category and indicated for diagnosis
Central Nervous System (CNS) Agents:	All products are covered without a PA <u>LENGTH OF AUTHORIZATIONS</u> : 365 Days
Fibromyalgia Agents	 NON PREFERRED CRITERIA: Must have had an inadequate clinical response of at least 14 days with at least two preferred drugs in different classes (see Additional Information section below) ADDITIONAL INFORMATION
	 Drugs and drug classes include gabapentin, pregabalin, short- and/or long-acting opioids, skeletal muscle relaxants, SNRIs, SSRIs, trazodone and tricyclic antidepressants
Central Nervous System (CNS)	BUPRENORPHINE SAFETY EDITS AND DRUG UTILIZATION REVIEW CRITERIA:



Agents: Medication Assisted Treatment of Opioid Addiction

• buprenorphine injection (SUBLOCADE) dosired to 300mg/30 days	18 solicatio will be limite
UPRENORPHINE INITIAL AUTHORIZATION CRITERIA	
afety edits are in place for dosages over 24mg of buprenorphine equivalents/day Pursuant to C osages exceeding 32mg of buprenorphine equivalents per day will not be approved.	hio Administrative Code 4731-33-03,
Has prescriber reviewed the OARRS within 7 days prior to the prior authorization request?	<u>Yes</u> No
Diagnosis (not approvable for pain) ICD-10 Code	
las individual been referred to counseling for addiction treatment?	Yes No N/A
f clinically indicated, has individual been offered a referral to counseling for addiction treatmen	?
Has the individual been offered a prescription for a naloxone kit?	<u>Yes</u> No
When is the individual's next appointment to assess induction therapy?	Date:
UPRENORPHINE RENEWAL CRITERIA Please provide the current duration of treatment as of the date of this request	
Please indicate the frequency of prescriber meetings.	
Has individual been actively participating in counseling AND been compliant with all	
sessions? Date of last counseling	<u>Yes</u> No
Has the dose been reduced in the past 6 months?	YesNo
Has there been an evaluation for a dose reduction since the previous PA request? If NO, please provide explanation	<u>YesNo</u>
Prescriber attests to the continued monitoring for safety and efficacy of the requested medication	_ <u>Yes</u> _No
Has the prescriber reviewed Ohio Automated Rx Reporting System within 7 days prior to the request?	PA <u>Yes</u> No
If individual is receiving opioids, benzodiazepines, sedative hypnotics, carisoprodol or tramadol, has the physician coordinated with all prescribers of controlled substances and determined treatment should continue?	<u>YesNo</u>
If YES, has an addiction specialist recommended to continue substance abuse treatment?	— Yes — No
Addiction Specialist consulted Phone Number	-Date
Is the individual receiving opioids, benzodiazepines, sedative hypnotics, carisoprodol or tramadol?	YesNo
Skip this question if answering "No" to the previous question	Yes No
Has the prescriber coordinated care with the prescriber(s) of the above listed substances and	
evaluated the risks and benefits of the combined use of these medications? Lab testing requirements met (at least twice per guarter for first year of treatment, once per-	
quarter thereafter)?	<u>— Yes — No</u>



REQUIRED FOR ALL BUPRENORPHINE REQUESTS	
ls the individual pregnant?	<u>YesNo</u>
Is the individual breastfeeding?	<u>YesNo</u>
Has the individual been explained the difference between an allergic reaction and symptoms of opioid withdrawal?	Yes No
Does the patient have an allergy or other contraindication to Naloxone? Please list reactions or reasons for contraindications	_ <u>Yes</u> Ne
Does the patient have an allergy to naloxone? If yes, please select appropriate box and provide additional information if needed.	Yes No Anaphylaxis Bronchospasm
	Angioneurotic edemaAnaphylactic shockHives Swelling of face or mouth Other:
Does the patient have a contraindication to naloxone? If yes, please select appropriate box and provide additional information if needed.	YesNoPregnancyBreastfeedingModerate to severe hepatic impairment as evidenced
	by Child-Pugh Class B or COther:
Skip this question if the patient has an allergy or contraindication to naloxone	Methadone
Is the patient starting buprenorphine induction following use of methadone, fentanyl, or extended-release opioids? If yes, please select appropriate box and provide additional information if needed.*	Fentanyl Extended-release opioids
*This rationale can only be used for a 30-day authorization.	Yes N/A
Skip this question if the patient has an allergy or contraindication to naloxone	IesIV/A
is the patient going to covert from buprenorphine mono-product to buprenorphine-naloxone combination? If yes, please provide additional information if needed.*	
*This rationale can only be used for a one-time 30-day authorization. Additional Information	
Additional montageon	



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	REQUIRED FOR ALL LOFEXIDINE (LUCEMYRA) REQUESTS
	Provide medical justification supporting why an opioid taper (such as buprenorphine or methadone) cannot be used:
	Provide documentation of an inadequate clinical response (including trial dates) or contraindication to clonidine:
	To be exempt from the above lofexidine criteria, provide documentation that the drug was initiated in an inpatient setting:
	"Lucemyra length of authorization is 14 days
	SIGNATURE AND DATE
	attest that I am a member of the prescriber's staff in accordance with Ohio Administrative Code rule
	5160-9-03, as applicable. Only the prescribing provider or a member of the prescribing provider's staff may request prior authorization.
	Prescriber's Signature (or staff of prescriber) Date
	IF a staff member is attesting, please print your name
Central Nervous	NON-PREFERRED CRITERIA:
System (CNS)	Must have had an inadequate clinical response of at least 30 days
Agents: Multiple	with at least <u>one preferred</u> drug in this UPDL category and indicate
Sclerosis* LEGACY	for diagnosis
CATEGORY	
Central Nervous	NON-PREFERRED CRITERIA:
System (CNS)	Must have had an inadequate clinical response with at least two
Agents: Narcolepsy	preferred drugs - either at least 30 days of armodafinil or modafinil;
	OR at least 7 days of a preferred amphetamine or methylphenidate
	drug in this UPDL category and indicated for diagnosis
Central Nervous	NON-PREFERRED CRITERIA:
System (CNS)	Must have had an inadequate clinical response of at least 30 days
Agents: Neuropathic	with at least two preferred drugs within the same sub-section
Pain	classification in different drug classes in this UPDL category and
	indicated for diagnosis
	<u>ADDITIONAL GABAPE<mark>NTIN (GRALISE) AND GABAPENTIN ENCARBIL</mark></u>
	(HORIZANT) CRITERIA
	 Must have had an inadequate clinical response to a preferred
	gabapentin product



Central Nervous	NON-PREFERRED CRITERIA:
System (CNS)	Must have had an inadequate clinical response of at least <u>30 days</u>
Agents: Parkinson's	with at least one preferred drug within the same mechanism of
Agents	action sub-section classification in this UPDL category, if and
	indicated for diagnosis. , if available
Central Nervous	NON-PREFERRED CRITERIA:
System (CNS)	Must have had an inadequate clinical response of at least 30 days
Agents: Restless	with at least one preferred drug in this UPDL category and indicated
Legs Syndrome	for diagnosis
2085 0 7 11 11 11 11	To a diagnosis
Central Nervous	NON-PREFERRED CRITERIA:
System (CNS)	 Must have had an inadequate clinical response of at least 7 days
Agents: Sedative-	with at least two preferred drugs in this UPDL category and
Hypnotics, Non-	indicated for diagnosis
Barbiturate	
Central Nervous	NON-PREFERRED CRITERIA:
System (CNS)	Must have had an inadequate clinical response of at least 30 days
Agents: Skeletal	with at least two preferred drugs in this UPDL category and
Muscle Relaxants,	indicated for diagnosis
Non-	
Benzodiazepine	
Dermatologic	NON-PREFERRED CRITERIA:
Agents: Oral Acne	Must have had an inadequate clinical response of at least <u>90 days</u>
Products	with at least two preferred drugs in this UPDL category and
Troducts	indicated for diagnosis
Dermatologic	NON-PREFERRED CRITERIA:
Agents: Topical	Must have had an inadequate clinical response with at least two
Acne Products	preferred drugs within the same mechanism of action sub-section
Actie i Toddets	classification in this UPDL category. Trials must be 30 days for
	preferred non-retinoids and 90 days for preferred retinoids.
Endocrine Agents:	NON-PREFERRED CRITERIA:
Androgens	Must have had an inadequate clinical response of at least <u>90 days</u>
Androgens	with <u>ALL preferred</u> drugs in this UPDL category and indicated for
	diagnosis
Endocrine Agents:	NON-PREFERRED CRITERIA:
Diabetes –	
Hypoglycemia	 Must have had an inadequate clinical response of at least <u>one</u> <u>preferred</u> drug in this UPDL category <u>and indicated for diagnosis</u> OR
Treatments	the inability of the member and/or caregiver to administer a
i i catiliciits	,
Endocrino Aconto	preferred glucagon product in a timely fashion
Endocrine Agents: Diabetes – Insulin	NON-PREFERRED CRITERIA:
Diabetes – Insulin	Must have had an inadequate clinical response (defined as the inability to reach target A1C) after at least 120 days with at least
	inability to reach target A1C) after at least 120 days with at least
	two preferred drugs having a similar duration of action in this UPDL
Find and the Arms	category and indicated for diagnosis, if available
Endocrine Agents:	NON-PREFERRED CRITERIA:
Diabetes – Non-	Must have had an inadequate clinical response of at least 120 days
Insulin	with at least three preferred drugs in this UPDL category and
	indicated for diagnosis, if available if available.
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Endocrine Agents:	NON-PREFERRED CRITERIA:
Endometriosis	 Must have had an inadequate clinical response of at least <u>84 days</u>
	with at least one preferred step-therapy drug in this UPDL category
	and indicated for diagnosis
Endocrine Agents:	NON-PREFERRED CRITERIA:
Estrogenic Agents	Must have had an inadequate clinical response of at least 30 days
	with at least two preferred drugs in this UPDL category within the
	same route of administration, sub-section classification and if
	indicated for diagnosis, if available.
Endocrine Agents:	NON-PREFERRED CRITERIA:
Growth Hormone	Must have had an inadequate clinical response of at least <u>90 days</u>
	with at least one preferred drug within the same sub-section
	classification in this UPDL category and indicated for diagnosis of
	similar duration of action in this UPDL category
Endocrine Agents:	NON-PREFERRED CRITERIA:
Osteoporosis – Bone	Must have had an inadequate clinical response of at least <u>365 days</u>
Ossification	with at least one preferred drug within the same sub-section
Enhancers	classification in this UPDL category and indicated for diagnosis with
Limancers	the same mechanism of action if available
Gastrointestinal	NON-PREFERRED CRITERIA:
Agents: Anti-	
Emetics	Must have had an inadequate clinical response of at least 3 days with at least and professed drug in this LIDDL extension.
Linetics	at least one preferred drug in this UPDL category within the same
	mechanism of action, if sub-section classification and indicated for
Gastrointestinal	diagnosis , if available .
	NON-PREFERRED CRITERIA:
Agents: Bowel	Must have had an inadequate clinical response with at least <u>one</u> professed drug in this LIDDL category and indicated for diagnosis.
Preparations	preferred drug in this UPDL category and indicated for diagnosis
Gastrointestinal	NON-PREFERRED CRITERIA:
Agents: Crohn's	Must have had an inadequate clinical response of at least 30 days with at least two graftered draws in this URBL category.
Disease	with at least two preferred drugs in this UPDL category and
0	indicated for diagnosis
Gastrointestinal	NON-PREFERRED CRITERIA:
Agents: Irritable	Must have had an inadequate clinical response of at least 14 days with at least are professed days and are stored by a second days.
Bowel Syndrome	with at least one preferred drug and one step therapy drug in this
(IBS) with Diarrhea	UPDL category and indicated for diagnosis
Gastrointestinal	NON-PREFERRED CRITERIA:
Agents: Pancreatic	Must have had an inadequate clinical response of at least 14 days With at least two professed days in this URBL patents. The state of the sta
Enzymes	with at least two preferred drugs in this UPDL category and indicated
	for diagnosis
Gastrointestinal	NON-PREFERRED CRITERIA:
Agents: Proton	Must have had an inadequate clinical response of at least 30 days
Pump Inhibitors	with at least two preferred drugs in this UPDL category and
	indicated for diagnosis
Gastrointestinal	NON-PREFERRED CRITERIA:
Agents: Ulcerative	Must have had an inadequate clinical response of at least 30 days
Colitis	with at least two preferred drugs in this UPDL category within the
	same route of administration, if sub-section classification and
	indicated for diagnosis, if available



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Gastrointestinal	NON-PREFERRED CRITERIA:
Agents: Unspecified	Must have had an inadequate clinical response of at least 14 days
GI	with one step therapy drug this UPDL category and indicated for
5.	diagnosis , if indicated for diagnosis
Canitanninan	
Genitourinary	NON-PREFERRED CRITERIA:
Agents: Benign	Must have had an inadequate clinical response of at least <u>60 days</u>
Prostatic	with at least two preferred drugs, with at least one preferred within
Hyperplasia	the same sub-section classification with the same mechanism of
	action, if available and indicated for diagnosis, if available
Genitourinary	NON-PREFERRED CRITERIA:
Agents: Electrolyte	Must have had an inadequate clinical response of at least 14 days
Depleter Agents	with at least two one preferred step therapy drugs in this UPDL
Depicter Agents	category and indicated for diagnosis, if available in this UPDL
	category, one of which must have the same mechanism of action as
	the requested non-preferred drug, if available
Genitourinary	NON-PREFERRED CRITERIA:
Agents: Urinary	<ul> <li>Must have had an inadequate clinical response of at least <u>30 days</u></li> </ul>
Antispasmodics	with at least two preferred drugs in this UPDL category and
	indicated for diagnosis, one of which must be within the same sub-
	section classification, with different active ingredients within the
	same mechanism of action, if available
Hyperkalemia	NON-PREFERRED CRITERIA:
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Agents: Potassium	Must have had an inadequate clinical response of at least <u>30 days</u>
Binders	with at least <u>one preferred</u> drug in this UPDL category <mark>and indicated</mark>
	for diagnosis
Immunomodulator	NON-PREFERRED CRITERIA:
Immunomodulator Agents: Systemic	<ul> <li>NON-PREFERRED CRITERIA:</li> <li>Must have had an inadequate clinical response of at least <u>90 days</u></li> </ul>
Agents: Systemic	<ul> <li>Must have had an inadequate clinical response of at least <u>90 days</u> with at least <u>two preferred</u> drugs in this UPDL category that are not</li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least <u>90 days</u> with at least <u>two preferred</u> drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for</li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least <u>90 days</u> with at least <u>two preferred</u> drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis., if available</li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least <u>90 days</u> with at least <u>two preferred</u> drugs in this UPDL category that are not biosimilars of the same reference product, <u>if and indicated for diagnosis</u>. <u>if available</u></li> <li>For non-preferred <u>biosimilars</u> immunomodulators: must</li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least <u>90 days</u> with at least <u>two preferred</u> drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis., if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to</li> </ul> </li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least <u>90 days</u> with at least <u>two preferred</u> drugs in this UPDL category that are not biosimilars of the same reference product, <u>if and</u> indicated for diagnosis., <u>if available</u> <ul> <li>For non-preferred <del>biosimilars</del> immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL</li> </ul> </li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least <u>90 days</u> with at least <u>two preferred</u> drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis., if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to</li> </ul> </li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least <u>90 days</u> with at least <u>two preferred</u> drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis., if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available</li> </ul> </li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least 90 days with at least two preferred drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis., if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available</li> </ul> </li> <li>ADDITIONAL PRURIGO NODULARIS CRITERIA:</li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least <u>90 days</u> with at least <u>two preferred</u> drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis., if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available</li> </ul> </li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least 90 days with at least two preferred drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis., if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available</li> </ul> </li> <li>ADDITIONAL PRURIGO NODULARIS CRITERIA:</li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least 90 days with at least two preferred drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis, if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available</li> </ul> </li> <li>ADDITIONAL PRURIGO NODULARIS CRITERIA:         <ul> <li>Must be prescribed by or in consultation with a specialist (i.e.,</li> </ul> </li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least 90 days with at least two preferred drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis, if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available</li> </ul> </li> <li>ADDITIONAL PRURIGO NODULARIS CRITERIA:         <ul> <li>Must be prescribed by or in consultation with a specialist (i.e.,</li> </ul> </li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least 90 days with at least two preferred drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis., if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available</li> </ul> </li> <li>ADDITIONAL PRURIGO NODULARIS CRITERIA:         <ul> <li>Must be prescribed by or in consultation with a specialist (i.e., dermatologist, rheumatologist)</li> </ul> </li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least 90 days with at least two preferred drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis, if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available</li> </ul> </li> <li>ADDITIONAL PRURIGO NODULARIS CRITERIA:         <ul> <li>Must be prescribed by or in consultation with a specialist (i.e., dermatologist, rheumatologist)</li> </ul> </li> <li>Must provide documentation of an inadequate clinical response of</li> </ul>
Agents: Systemic Inflammatory	<ul> <li>Must have had an inadequate clinical response of at least 90 days with at least two preferred drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis, if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available</li> </ul> </li> <li>ADDITIONAL PRURIGO NODULARIS CRITERIA:         <ul> <li>Must be prescribed by or in consultation with a specialist (i.e., dermatologist, rheumatologist)</li> </ul> </li> <li>Must provide documentation of an inadequate clinical response of</li> </ul>
Agents: Systemic Inflammatory Disease  Infectious Disease	Must have had an inadequate clinical response of at least 90 days with at least two preferred drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis, if available  For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available  ADDITIONAL PRURIGO NODULARIS CRITERIA:  Must be prescribed by or in consultation with a specialist (i.e., dermatologist, rheumatologist)  Must provide documentation of an inadequate clinical response of at least 90 days with a topical steroid  NON-PREFERRED CRITERIA:
Agents: Systemic Inflammatory Disease  Infectious Disease Agents: Antibiotics –	<ul> <li>Must have had an inadequate clinical response of at least 90 days with at least two preferred drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis, if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available</li> </ul> </li> <li>ADDITIONAL PRURIGO NODULARIS CRITERIA:         <ul> <li>Must be prescribed by or in consultation with a specialist (i.e., dermatologist, rheumatologist)</li> </ul> </li> <li>Must provide documentation of an inadequate clinical response of at least 90 days with a topical steroid</li> <li>NON-PREFERRED CRITERIA:         <ul> <li>Must have had an inadequate clinical response of at least 3 days</li> </ul> </li> </ul>
Agents: Systemic Inflammatory Disease  Infectious Disease	<ul> <li>Must have had an inadequate clinical response of at least 90 days with at least two preferred drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis, if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available</li> </ul> </li> <li>ADDITIONAL PRURIGO NODULARIS CRITERIA:         <ul> <li>Must be prescribed by or in consultation with a specialist (i.e., dermatologist, rheumatologist)</li> </ul> </li> <li>Must provide documentation of an inadequate clinical response of at least 90 days with a topical steroid</li> <li>NON-PREFERRED CRITERIA:         <ul> <li>Must have had an inadequate clinical response of at least 3 days with at least one preferred drug in this UPDL category and indicated</li> </ul> </li> </ul>
Agents: Systemic Inflammatory Disease  Infectious Disease Agents: Antibiotics – Cephalosporins	<ul> <li>Must have had an inadequate clinical response of at least 90 days with at least two preferred drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis, if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available</li> </ul> </li> <li>ADDITIONAL PRURIGO NODULARIS CRITERIA:         <ul> <li>Must be prescribed by or in consultation with a specialist (i.e., dermatologist, rheumatologist)</li> </ul> </li> <li>Must provide documentation of an inadequate clinical response of at least 90 days with a topical steroid</li> <li>NON-PREFERRED CRITERIA:         <ul> <li>Must have had an inadequate clinical response of at least 3 days with at least one preferred drug in this UPDL category and indicated for diagnosis</li> </ul> </li> </ul>
Agents: Systemic Inflammatory Disease  Infectious Disease Agents: Antibiotics – Cephalosporins  Infectious Disease	<ul> <li>Must have had an inadequate clinical response of at least 90 days with at least two preferred drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis, if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available</li> </ul> </li> <li>ADDITIONAL PRURIGO NODULARIS CRITERIA:         <ul> <li>Must be prescribed by or in consultation with a specialist (i.e., dermatologist, rheumatologist)</li> </ul> </li> <li>Must provide documentation of an inadequate clinical response of at least 90 days with a topical steroid</li> <li>NON-PREFERRED CRITERIA:         <ul> <li>Must have had an inadequate clinical response of at least 3 days with at least one preferred drug in this UPDL category and indicated for diagnosis</li> </ul> </li> <li>NON-PREFERRED CRITERIA:</li> <li>NON-PREFERRED CRITERIA:</li> </ul>
Agents: Systemic Inflammatory Disease  Infectious Disease Agents: Antibiotics – Cephalosporins	<ul> <li>Must have had an inadequate clinical response of at least 90 days with at least two preferred drugs in this UPDL category that are not biosimilars of the same reference product, if and indicated for diagnosis, if available         <ul> <li>For non-preferred biosimilars immunomodulators: must provide documentation of inadequate clinical response to its preferred reference product or biosimilar, in this UPDL category, if and indicated for the diagnosis, if available</li> </ul> </li> <li>ADDITIONAL PRURIGO NODULARIS CRITERIA:         <ul> <li>Must be prescribed by or in consultation with a specialist (i.e., dermatologist, rheumatologist)</li> </ul> </li> <li>Must provide documentation of an inadequate clinical response of at least 90 days with a topical steroid</li> <li>NON-PREFERRED CRITERIA:         <ul> <li>Must have had an inadequate clinical response of at least 3 days with at least one preferred drug in this UPDL category and indicated for diagnosis</li> </ul> </li> </ul>



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	for diagnosis
Infectious Disease	NON-PREFERRED CRITERIA:
Agents: Antibiotics –	 Must have had an inadequate clinical response of at least <u>3 days</u>
Macrolides	with at least one preferred drug in this UPDL category and indicated
	for diagnosis
Infectious Disease	NON-PREFERRED CRITERIA:
Agents: Antibiotics –	Must have had an inadequate clinical response of at least 3 days
Quinolones	with at least one preferred drug in this UPDL category and indicated
	for diagnosis
Infectious Disease	NON-PREFERRED CRITERIA:
Agents: Antibiotics –	 Must have had an inadequate clinical response of at least 3 days
Tetracyclines	with at least one preferred drug for acute infections OR at least 90
	days with at least one preferred oral drug for acne in this UPDL
	category and indicated for diagnosis
Infectious Disease	NON-PREFERRED CRITERIA:
Agents: Antifungals	 Must have had an inadequate clinical response of at least 3 days
	with at least <u>two preferred</u> drugs , if indicated for the diagnosis in
	this UPDL category in this UPDL category and indicated for diagnosis
Infectious Disease	NON-PREFERRED CRITERIA:
Agents: Antivirals –	 Must have had an inadequate clinical response defined as not
Hepatitis C Agents	achieving sustained virologic response (SVR) with guideline-
	recommended preferred drugs in this UPDL category and indicated
	for diagnosis
Infectious Disease	NON-PREFERRED CRITERIA:
Agents: Antivirals –	 Must have had an inadequate clinical response of at least <u>3 days</u>
Herpes	with at least <u>one preferred</u> drug in this UPDL category <mark>and indicated</mark>
	for diagnosis
Infectious Disease	FOSTEMSAVIR (RUKOBIA) CRITERIA:
Agents: Antivirals –	 Must provide documentation of a multidrug-resistant HIV-1
HIV* LEGACY	<mark>infection</mark>
CATEGORY	
	NON-PREFERRED CRITERIA:
	Must have had an inadequate clinical response of at least <u>30 days</u> with at least an appropriate days in this URBN acts are a limited at least and the standard of the s
	with at least one preferred drug in this UPDL category and indicated
	for diagnosis. If applicable, the request must address the inability to
Onbtholmic Access	use the individual components.
Ophthalmic Agents:	NON-PREFERRED CRITERIA:
Antibiotic and	Must have had an inadequate clinical response of at least <u>3 days</u> with at least two professed days in this LIDBL coto says and
Antibiotic-Steroid	with at least two preferred drugs in this UPDL category and
Combination Drops and Ointments	indicated for diagnosis
	NON DEFERDED CRITERIA
Ophthalmic Agents: Antihistamines &	 NON-PREFERRED CRITERIA: Must have had an inadequate clinical response of at least 7 days
Mast Cell Stabilizers	· · · · · · · · · · · · · · · · · · ·
iviasi Celi Stabilizeis	with at least <u>two preferred</u> drugs in this UPDL category <mark>and</mark>
	indicated for diagnosis
Onhthalmic Agents:	indicated for diagnosis STED THERAPY CRITERIA:
Ophthalmic Agents: Dry Eye Treatments	 indicated for diagnosis STEP THERAPY CRITERIA: Must have had an inadequate clinical response of at least 14 days



	with at least <u>one</u> preferred drug in this <mark>UPDL</mark> category in the previous 120 days
	NON-PREFERRED CRITERIA:
	Must have had an inadequate clinical response of at least 14 days
	with at least two preferred drugs in this UPDL category and
	indicated for diagnosis
Ophthalmic Agents:	STEP THERAPY CRITERIA:
Glaucoma Agents	Must have had an inadequate clinical response of at least <u>30 days</u>
	with at least one preferred drug in the same sub-section
	classification in this UPDL category and indicated for diagnosis, if
	available
	NON-PREFERRED CRITERIA:
	Must have had an inadequate clinical response of at least 30 days
	with at least two preferred drugs within the same sub-section
	classification in this UPDL category and indicated for diagnosis in
	the same class, if available
Ophthalmic Agents:	NON-PREFERRED CRITERIA:
NSAIDs	 Must have had an inadequate clinical response of at least <u>3 days</u>
	with at least two preferred drugs in this UPDL category and
	indicated for diagnosis
Ophthalmic Agents:	NON-PREFERRED CRITERIA:
Ophthalmic Steroids	Must have had an inadequate clinical response of at least <u>7 days</u> with at least two preferred drugs in this URDI estagery and
	with at least <u>two preferred</u> drugs in this UPDL category <mark>and indicated for diagnosis</mark>
Otic Agents:	NON-PREFERRED CRITERIA:
Antibacterial and	Must have had an inadequate clinical response of at least 3 days
Antibacterial/Steroi	with at least <u>two preferred</u> drugs in this UPDL category <mark>and</mark>
d Combinations	indicated for diagnosis
Respiratory Agents:	NON-PREFERRED CRITERIA:
Antihistamines –	Must have had an inadequate clinical response of at least 7 days
Second Generation	with at least two different preferred drugs in this UPDL category
	and indicated for diagnosis .
Respiratory Agents:	NON-PREFERRED CRITERIA:
Cystic Fibrosis	Must have had an inadequate clinical response of at least <u>30 days</u>
	with at least one preferred drug in this UPDL category and indicated
	for diagnosis
Respiratory Agents:	NON-PREFERRED CRITERIA:
Epinephrine	Must have had an inadequate clinical response to at least <u>one</u> professed drug in this URDL category and indicated for diagnosis.
Respiratory Agents:	preferred drug in this UPDL category and indicated for diagnosis NON-PREFERRED CRITERIA:
Hereditary	Must have had an inadequate clinical response of at least 3 days
Angioedema	with at least one preferred acute drug in this UPDL category and
	indicated for diagnosis to request a non-preferred acute drug.
	Must have had an inadequate clinical response of at least <u>14 days</u>
	with at least one preferred prophylaxis drug in this UPDL category



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	and indicated for diagnosis to request a non-preferred prophylaxis
	drug.
Respiratory Agents:	NON-PREFERRED CRITERIA:
Inhaled Agents	 Must have had an inadequate clinical response of at least 14 days with at least two preferred drugs in this UPDL category within the same mechanism of action, if sub-section classification and indicated for diagnosis, if available.
Respiratory Agents:	NON-PREFERRED CRITERIA:
Leukotriene	Must have had an inadequate clinical response of at least <u>30 days</u>
Receptor Modifiers	with at least two preferred drugs in this UPDL category and
& Inhibitors	indicated for diagnosis
Respiratory Agents:	CLINICAL PA CRITERIA:
Monoclonal	• For Chronic Obstructive Pulmonary Disease (COPD):
Antibodies-Anti-	For Chronic Obstructive Pulmonary Disease (COPD).
IL/Anti-IgE	 The patient must have an eosinophilic count of greater
IL/AIIU-IgE	than or equal to 300 cells per mcL within 12 months prior
	to initiation of therapy AND
	 The patient has a history of uncontrolled disease, as
	indicated by greater than or equal to 2 COPD
	exacerbations or greater than or equal to 1 COPD
	exacerbation resulting in a hospitalization despite being on
	standard of care, defined as triple therapy
	(LAMA+LABA+ICS) for at least 3 months prior to request,
	and at a stable dose for at least 1 month prior.
	NON-PREFERRED CRITERIA:
	Must have had an inadequate clinical response of at least <u>90 days</u>
	with at least one preferred drug in this UPDL category and indicated
	for diagnosis
	- Constant of the constant of
Respiratory Agents:	NON-PREFERRED CRITERIA:
Nasal Preparations	 Must have had an inadequate clinical response of at least 14 days
	with at least two preferred drugs in this UPDL category within the
	same mechanism of action, if sub-section classification and
	indicated for diagnosis. , if available
Respiratory Agents:	NON-PREFERRED CRITERIA:
Pulmonary Fibrosis	 Must have had an inadequate clinical response of at least <u>30 days</u>
	with at least <u>one preferred</u> drug in this UPDL category <mark>and indicated</mark>
	for diagnosis
Topical Agents:	NON-PREFERRED CRITERIA:
Antifungals	 Must have had an inadequate clinical response of at least 14 days
	with at least two preferred drugs in this UPDL category and
	indicated for diagnosis , if indicated for diagnosis
Topical Agents:	NON-PREFERRED CRITERIA:
Antiparasitics	Must have had an inadequate clinical response of at least 14 days
	with at least one preferred drug in this UPDL category and indicated
	for diagnosis



Topical Agents:	NON-PREFERRED CRITERIA:
Corticosteroids	Must have had an inadequate clinical response of at least <u>14 days</u>
	with at least two preferred drugs within the same sub-section
	classification in this UPDL category and indicated for diagnosis, if
	<mark>available</mark> of similar potency
Topical Agents:	NON-PREFERRED CRITERIA:
Immunomodulators	Must have had an inadequate clinical response of at least 30 days
	with at least one preferred drug in this UPDL category and indicate
	for diagnosis

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