

Formulary Exception/Prior Authorization Request Form

Patient Information Patient Name:			Prescriber Information Prescriber Name:			
Patient ID#:						
Address:		Address:				
City:	State:	City:			State:	
Home Phone:	ZIP:	Office Pho	ne #:	Office Fax #:	ZIP:	
Condon Mon F	DOB:	Contact Po	erson at Doo	ctor's Office:		
Gender: M or F						
Medication:	Diagnosis and Strength:	Medical Info	rmation	Directions for use (F	Frequency):	
Expected Length of Therapy:	Qty:	Day Supply:		a continuation of therapy		
Diagnosis:		Diagnosis	(ICD) Code	(s):		
	LL RELEVANT CLINICAL DOC slisted on page 2. For any drugs					
Expedited/Urgent Review Reque frame may seriously jeopardize the						
ase list all medications the patient has tried	·	•		,		
Medication name, reason for failure, include						
Drug(s) contraindicated:						
Adverse event (e.g., toxicity, allergy) for ea						
	onic conditions (e.g., psychiatric	condition, ep	ilepsy, dem	entia) who is stable on th	ne current drug(s) and who might	
gh risk for a significant adverse event with						
	rmed by diagnostic testing? <i>If ye</i>	es, please pro	ovide diagn	ostic test and date:		
igh risk for a significant adverse event with a			_			

Is the request for Diabetic Test Strips? If yes, please answer the two questions below.

1. Does the patient have an insulin pump? If so, please provide make and model (e.g., OmniPod, MiniMed 530G)

Does the patient have an insulin pump that is incompatible with Accu-Chek products? Yes or No

PRESCRIPTION BENEFIT PLAN MAY REQUEST ADDITIONAL INFORMATION OR CLARIFICATION, IF NEEDED, TO EVALUATE REQUESTS. FOR NON-SPECIALTY DRUGS, PLEASE FAX COMPLETED FORM TO 1-844-814-2258.

FOR SPECIALTY DRUGS, PLEASE FAX COMPLETED FORM TO 1-844-814-2259.

in m	attest that the medication requested is medically necessary for this patient. I further attest that the information provided is accurate and true, and that documentation supporting this formation is available for review if requested by Oscar or, if applicable, a state or federal regulatory agency. I understand that any person who knowingly makes or causes to be ade a false record or statement that is material to a claim ultimately paid by the United States government or any state government may be subject to civil penalties and treble amages under both the federal and state False Claims Acts. See, e.g., 31 U.S.C. §§ 3729-3733.
D	Date:
Г	rescriber Signature:
he	onfidentiality Notice: The documents accompanying this transmission contain confidential health information that is legally privileged. If you are not the intended recipient, you are ereby notified that any disclosure, copying, distribution of these documents is strictly prohibited. If you have received this information in error, please notify the sender immediately ia return fax) and arrange for the return or destruction of these documents.
	ASE COMPLETE CORRESPONDING SECTION FOR THESE SPECIFIC DRUGS/CLASSES LISTED BELOW AND CIRCLE THE APPROPRIATE WER OR SUPPLY RESPONSE.
」 1.	ANTIFUNGALS: Is the request for terbinafine (Lamisil), Kerydin or Jublia? (please circle one)
2.	Does the patient have a diagnosis of onychomycosis due to tinea unguium, Trichophyton rubrum or Trichophyton mentagrophytes? Yes or No (circle
	appropriate diagnosis) If yes to question 2, is the onychomycosis confirmed by a fungal diagnostic test? Yes or No
3. 4.	Does the infection involve the toenails, fingernails or both? (please circle) Is the request for treatment of tinea corporis or tinea cruris in a patient who is immunocompromised or has extensive or complicated infection? Yes or No
	If yes to question 4, does the patient require systemic therapy or have more extensive superficial infections? Yes or No
5.	Has the patient experienced an inadequate treatment response, intolerance or contraindication to an oral antifungal therapy? Yes or No
] 1. 2.	ANTIEMETIC (5-HT3) AGENTS: Is the patient receiving moderate to highly emetogenic chemotherapy or receiving radiation therapy? Yes or No Is the patient pregnant with the diagnosis of Hyperemesis Gravidarum and a documented risk for hospitalization? Yes or No If yes to question 2, has the patient experienced an inadequate treatment response, intolerance or contraindication to two of the following medications: Vitamin B6, doxylamine, doxylamine/pyridoxine extended-release (Boniesta), doxylamine/pyridoxine delayed-release (Diclegis), promethazine (Phenergan), trimethobenzamide (Tigan) or metoclopramide (Reglan)? Yes or No (if yes, circle appropriate medications)
7	ERECTILE DYSFUNCTION:
1. 2.	Is the drug being prescribed for erectile dysfunction? Yes or No Is the drug being prescribed for symptomatic Benign Prostatic Hyperplasia (BPH)? Yes or No
]	INSOMNIA AGENTS:
1. 2.	Does the patient have a diagnosis of insomnia? Yes or No Have potential causes of sleep disturbances been addressed (e.g., inappropriate sleep hygiene and sleep environment issues, treatable medical/psychological causes of chronic insomnia)? Yes or No
]	PROTON PUMP INHIBITORS:
1.	Does the patient have endoscopically verified peptic ulcer disease OR frequent and severe symptoms of gastroesophageal reflux disease (GERD) (e.g., heartburn, regurgitation) OR atypical symptoms or complications of GERD (e.g., dysphagia, hoarseness, erosive esophagitis)? Yes or No (if yes, please circle one)
2.	Does the patient have Barrett's esophagus as confirmed by biopsy OR a Hypersecretory syndrome (e.g. Zollinger-Ellison) confirmed with a diagnostic test?
3.	Yes or No (if yes, please circle one) Is the patient at high risk for GI adverse events? Yes or No
	PROVIGIL/NUVIGIL:
1. 2.	Does the patient have a diagnosis of Shift Work Disorder (SWD)? Yes or No Does the patient have a diagnosis of Obstructive Sleep Apnea confirmed by polysomnography? Yes or No
3.	Does the patient have a diagnosis of Narcolepsy confirmed by sleep lab evaluation? Yes or No
4.	Is the request for Provigil, and does the patient have a diagnosis of fatigue related to multiple sclerosis? Yes or No If yes to question 4, has the patient had an inadequate treatment response, intolerance or contraindication to amantadine? Yes or No
]	STIMULANTS: AMPHETAMINES, METHYLPHENIDATES, STRATTERA
1. 2.	Does the patient have a diagnosis of attention deficit/hyperactivity disorder (ADHD) or attention deficit disorder (ADD)? Yes or No Has the diagnosis been documented (i.e., complete clinical assessment, using DSM-5®, standardized rating scales, interviews/questionnaires)? Yes or No
3.	Does the patient have a diagnosis of Narcolepsy confirmed by sleep study? Yes or No
4.	Does the patient have a diagnosis of moderate to severe binge eating disorder (BED)? Yes or No
] 1.	TRETINOIN PRODUCTS: Does the patient have the diagnosis of acne vulgaris or keratosis follicularis (Darier's disease, Darier-White disease)? Yes or No (if yes, please circle one)
]	TAZORAC:
1. 2.	Does the patient have a diagnosis of acne vulgaris? Yes or No
2. 3. 4.	Does the patient have a diagnosis of plaque psoriasis? Yes or No Will the patient be applying Tazorac to less than 20 percent of body surface area? Yes or No Has the patient had intolerance, inadequate treatment response or contraindication to one topical corticosteroid? Yes or No
7	TESTOSTERONE PRODUCTS:
1.	Does the patient have primary or secondary (hypogonadotropic) hypogonadism? Yes or No
2. 3.	Does the patient have age-related hypogonadism? Yes or No Does the patient have at least two confirmed low testosterone levels according to current practice guidelines or your standard male lab reference values?
1	Yes or No
4.	Is the drug being prescribed for female-to-male gender reassignment? Yes or No

TRIPTANS: Does the patient have confirmed or suspected cardiovascular or cerebrovascular disease, or uncontrolled hypertension? Yes or No Does the patient have a diagnosis of migraine headache or cluster headache? Please circle one

Is the patient currently using or unable to use migraine prophylactic therapy (e.g., amitriptyline, propranolol, timolol)? Yes or No

Has medication overuse headache been considered and ruled out? Yes or No

Does the patient need an amount for treating more than eight headaches per month with a 5-HT1 agonist? Yes or No

- Does the patient have osteoarthritis pain in joints susceptible to topical treatment such as feet, ankles, knees, hands, wrist or elbow? Yes or No
- Is the treatment with the requested drug necessary due to intolerance or a contraindication to oral nonsteroidal anti-inflammatory (NSAID) drugs? Yes or No
- Does the patient require more than 1000 grams (10 tubes) per month? Yes or No