



**Medical Exception/  
Prior Authorization/Precertification\*  
Request for Prescription Medications**

Fax this form to: 1-877-269-9916  
OR

Submit your request online at:  
<https://navinet.navimedix.com/Main.asp>  
Visit [www.aetna.com/formulary](http://www.aetna.com/formulary) to access  
our Pharmacy Clinical Policy Bulletins.

For FASTEST service, call 1-855-240-0535, Monday-Friday, 8 a.m. to 6 p.m. Central Time

Patient Information		Prescriber Information	
Patient Name		Today's Date	
Patient Insurance ID Number		Physician Name	
Patient Address, City, State, ZIP		Physician Address	
Home Telephone		M.D. Office Telephone Number	
Gender <input type="checkbox"/> Male <input type="checkbox"/> Female	Patient Date of Birth	M.D. Office Fax Number	
Diagnosis and Medical Information			
Medication		Strength	Frequency
Expected Length of Therapy	Quantity	Day Supply	If this is a continuation of therapy, how long has the patient been on the medication?
<b>PLEASE CHECK ALL BOXES THAT APPLY:</b>			
Do you want a drug specific prior authorization criteria form faxed to your office? <input type="checkbox"/> Yes <input type="checkbox"/> No (If yes, no further questions are required).			
<input type="checkbox"/> What condition is the drug being prescribed for? ICD code _____ Diagnosis _____			
<input type="checkbox"/> Does the patient have a diagnosis of cancer? <input type="checkbox"/> Yes <input type="checkbox"/> No			
<input type="checkbox"/> Please list all medications the patient has tried specific to the diagnosis and specify below: Therapeutic failure, including length of therapy for each drug: _____ Drugs (s) contraindicated: _____ Adverse even (e.g., toxicity, allergy) for each drug: _____			
<input type="checkbox"/> Is the request for a patient with one or more chronic conditions (e.g., psychiatric condition, diabetes) who is stable on the current drug(s) and who might be at high risk for a significant adverse event with a medication change? If so, specify anticipated significant adverse event: _____			
<input type="checkbox"/> Has the condition been confirmed by diagnostic testing? If so, please provide diagnostic test and date: _____			
<input type="checkbox"/> Does the patient have a clinical condition for which other alternatives are not recommended based on published guidelines or clinical literature? If so, please provide documentation: _____			
<input type="checkbox"/> Does the patient require a specific dosage form (e.g., suspension, solution, injection)? If so, please provide dosage form: _____			
<input type="checkbox"/> Are additional risk factors (e.g., GI risk, cardiovascular risk, age) present? If so, please provide risk factors: _____			
<input type="checkbox"/> Other: Please provide additional relevant information: _____			
REQUIRED CLINICAL INFORMATION: PLEASE PROVIDE ALL RELEVANT CLINICAL DOCUMENTATION TO SUPPORT USE OF THIS MEDICATION. PLEASE COMPLETE CORRESPONDING SECTION ON BACK PAGE FOR THE SPECIFIC DRUG/CLASS LISTED BELOW. Antifungals/Antiemetic (5-HT3) Agents/Celebrex/Erectile Dysfunction Agents/Proton Pump Inhibitors/Protopic Provigil/Nuvigil/Stimulants/Tazorac/Tretinoin Products/Triptans			
**FOR ANY DRUG/CLASS NOT LISTED ON THE BACK PAGE, PLEASE ATTACH ADDITIONAL INFORMATION, BUT CANNOT EXCEED TWO PAGES**			
<b>PRESCRIPTION BENEFIT PLAN MAY REQUEST ADDITIONAL INFORMATION OR CLARIFICATION, IF NEEDED, TO EVALUATE REQUESTS</b>			
I 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 information is available for review if requested by the health plan sponsor, or, if applicable, a state or federal regulatory agency. I understand that any person who knowingly makes or causes to be made 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 damages under both the federal and state False Claims Acts. See, e.g., 31 U.S.C. §§ 3729-3733.			
Prescriber Signature			Date

**Confidentiality Notice:** The documents accompanying this transmission contain confidential health information that is legally privileged. If you are not the intended recipient, you are hereby 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 (via return FAX) and arrange for the return or destruction of these documents.

**PLEASE COMPLETE CORRESPONDING SECTION FOR THESE SPECIFIC DRUGS/CLASSES LISTED BELOW AND CIRCLE THE APPROPRIATE ANSWER OR SUPPLY RESPONSE.**

**ANTIFUNGALS: LAMISIL, SPORANOX, PENLAC, DIFLUCAN**

Does the patient have secondary medical risk factors? Please specify which risk factor(s): \_\_\_\_\_

If the patient has a diagnosis of Onychomycosis, does the infection involve the toenails, fingernails or both? **Please circle**

If the diagnosis is Tinea corporis or Tinea cruris, does the patient require systemic therapy or have more extensive superficial infections?  Yes  No

**ANTIEMETIC (5-HT3) AGENTS: (Ondansetron quantities of 12 or less per 30 days do not require a prior authorization)**

Is the patient receiving moderate to highly emetogenic chemotherapy? Monthly frequency \_\_\_\_\_  Yes  No

Is the patient receiving radiation therapy? Monthly frequency \_\_\_\_\_  Yes  No

If the patient has a diagnosis of Hyperemesis Gravidarum, has the patient experienced an inadequate treatment response to two of the following medications?

vitamin B6, doxylamine, promethazine (Phenergan), trimethobenzamide (Tigan) or metoclopramide (Reglan)?  Yes  No

**CELEBREX:**

Is the patient at risk for a severe NSAID-related gastrointestinal (GI) adverse event (e.g., NSAID associated gastric ulcer, GI bleed)?  Yes  No

**ERECTILE DYSFUNCTION: CIALIS, LEVITRA, VIAGRA, ALPROSTADIL**

Does the patient require nitrate therapy on a regular OR on an intermittent basis, or is the patient currently taking another ED medication?  Yes  No

If a diagnosis of erectile dysfunction, is it due to neurogenic etiology, vasculogenic etiology, psychogenic etiology or mixed etiology? **Please circle.**

Is it being used for symptomatic Benign Prostatic Hyperplasia (BPH)?  Yes  No

**PROTON PUMP INHIBITORS:**

Does the patient have frequent and severe symptoms of GERD (e.g., heartburn, regurgitation)?  Yes  No

Does the patient have atypical symptoms or complications of GERD (e.g., dysphagia, hoarseness, erosive esophagitis)?  Yes  No

**PROTOPIC:**

Has the patient had a therapeutic failure of a topical corticosteroid?  Yes  No

**PROVIGIL/NUVIGIL:**

If the patient has a diagnosis of Obstructive Sleep Apnea, is the patient currently using a continuous positive airway pressure (CPAP) machine or other device?  Yes  No

**STIMULANTS: AMPHETAMINES, METHYLPHENIDATES, STRATTERA**

Is this a renewal of therapy?  Yes  No

**TAZORAC/ TRETINOIN PRODUCTS:**

Has the patient tried and failed products from the following categories: Salicylic Acid Products OR Benzoyl Peroxide products?  Yes  No

**TRIPTANS:**

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