COVERAGE EXCEPTION

PRESCRIBER FAX FORM

Only the prescriber may complete this form. This form is for prospective, concurrent, and retrospective reviews. **Incomplete forms will be returned for additional information**. The following documentation is required for preauthorization consideration. For formulary information please visit www.myprime.com. What is the priority level of this request? ☐ Standard ☐ Date of service (if applicable): ☐ Urgent (NOTE: Urgent is defined as when the prescriber believes that waiting for a standard review could seriously harm the patient's life, health, or ability to regain maximum function.) PATIENT AND INSURANCE INFORMATION Today's Date: Patient Name (First): Last: DOB (mm/dd/yyyy): City, State, Zip: Patient Address: Patient Telephone: Member ID Number: Group Number: PRESCRIBER/CLINIC INFORMATION Prescriber NPI#: Prescriber Name: Specialty: Contact Name: Clinic Name: Clinic Address: City, State, Zip: Phone #: Secure Fax #: RENDERING/SERVICING PRESCRIBER INFORMATION (IF APPLICABLE) Prescriber Name: Prescriber NPI#: Specialty: Contact Name: Clinic Name: Clinic Address: Phone # City, State, Zip: Secure Fax #: PLEASE ATTACH ANY ADDITIONAL INFORMATION THAT SHOULD BE CONSIDERED WITH THIS REQUEST Patient's Diagnosis (ICD code and description): Patient's height: Patient's weight: Medication Requested: Strength: Dosing Schedule: Quantity per Month: For all requests: If yes, please explain risk: 2. Please list all reasons for selecting the requested medication, strength, dosing schedule, and quantity over alternatives (e.g., contraindications, allergies, history of adverse drug reactions to alternatives, lower dose has been tried, information supporting dose over FDA). Please list any other medications the patient will use in combination with the requested medication for treatment of this diagnosis. If yes, please provide supporting information: Please continue to the next page.

Patient Name (First):		Last:		M: DOB (mm/dd/yyyy):				
5.	Please list all medications the pa	ltient has previously tried and failed for treatment of	this diagnosis.	Please specify if	the patient			
	has tried brand-name products, generic products or over-the-counter products.							
		Date(s):		Date(s):				
				Date(s):				
		Date(s):		Date(s):				
6.	Please provide information indicating the cause of the patient's failure to any previously tried treatments for this diagnosis							
Foi	Aspirin Therapy:							
7.	Is the patient pregnant, at high risk of preeclampsia, and using the requested agent after 12 weeks							
	gestation?			Yes	☐ No			
Foi	Bowel Prep Therapy:							
8.	Will the requested agent be used for the preparation of colorectal cancer screening using fecal occult blood							
	testing, sigmoidoscopy, or colono	oscopy?		🗌 Yes	☐ No			
Foi	Breast Cancer Primary Prevent	ion Therapy:						
9.	Is the requested agent being requested	uested for the primary prevention of breast cancer?		Yes	☐ No			
10.	Is the patient female?			🗌 Yes	☐ No			
	If no, is the requested agen	t medically appropriate for the patient's sex?		Yes	☐ No			
	If yes, please explain:							
Foi	Contraceptive Agents:							
11.	Is the requested agent being use	d for contraception?		🗌 Yes	☐ No			
12.	Is the patient female?			🗌 Yes	☐ No			
	If no, is the requested agen	t medically appropriate for the patient's sex?		🗌 Yes	☐ No			
	If yes, please explain:							
Foi	Folic Acid Therapy:							
13.	Is the requested agent being use	d to support pregnancy?		🗌 Yes	□No			
14.	Is the patient female?			🗌 Yes	☐ No			
	If no, is the requested agen	t medically appropriate for the patient's sex?		🗌 Yes	☐ No			
	If yes, please explain:							
Foi	HIV Infection PrEP Therapy:							
15.	Is the requested agent being use	d for PrEP?		Yes	☐ No			
16.	Is the requested agent medically	necessary?		🗌 Yes	☐ No			
	If yes, please explain:							
17.	Is the requested PrEP agent any	of the following: tenofovir disoproxil fumarate and e	mtricitabine co	mbination				
	ingredient agent, tenofovir alafen	amide and emtricitabine combination ingredient age	ent, or caboteg	ravir? 🗌 Yes	☐ No			
18.	Does the patient have an increas	ed risk for HIV infection?		Yes	☐ No			
19.	Has the patient recently tested no	egative for HIV?		Yes	☐ No			
Ple	ase continue to the next page.							

Patient Name (First):	Last:	M:	DOB (mm/dd/yyyy):				
For Infant Eye Ointment Therapy:							
20. Is the requested agent for the prevention of gonococcal ophthalmia neonatorum?							
For Iron Supplements Therapy:							
21. Is the patient at increased risk of iron deficiency anemia?							
For Statin Therapy:							
22. Is the requested agent for use in the primary prevention of cardiovascular disease (CVD)?							
23. Does the patient have any of the following CVD risk factors? Check all that apply.							
☐ Dyslipidemia ☐	Hypertension						
☐ Diabetes ☐	Smoking						
24. Does the patient have a calculated 10-year risk of a cardiovascular event of 10% or greater based on							
calculations from the ACA/AHA ASCVD Risk Estimator (https://tools.acc.org/ASCVD-Risk-Estimator/)?							
For Tobacco Cessation Therapy:							
25. Is the patient a non-pregnant adult?			🗌 Yes 🔲 No				
26. Has the patient received 180 or more day supply of the requested tobacco cessation agent type (e.g.,							
NRT, bupropion, varenicline) in the past 365 days? Yes No							
If yes, is the patient currently being treated with the requested tobacco cessation agent type (e.g., NRT, bupropion,							
varenicline) and is expected to be succ	cessful on this course of therapy?		Yes No				
If yes, please explain:							
If no, is there support for the anticipated success of repeating therapy with the requested tobacco cessation agent							
type (e.g., NRT, bupropion, varenicline)? 🗌 Yes 🔲 No							
If yes, please provide supporting information:							
For Vaccine Therapy:							
27. Will the requested vaccine be used per the recommendations of the Advisory Committee on Immunization							
Practices (ACIP) and Centers for Disease Control (CDC)?							
Please fax or mail this form to:	CONFIDENTIALITY NOTICE:	This commun	ication is intended only for the				
Prime Therapeutics LLC Clinical Review Department	use of the individual entity to wl	use of the individual entity to which it is addressed and may contain					
2900 Ames Crossing Road Suite 200 Eagan, MN 55121	information that is privileged or	information that is privileged or confidential. If the reader of this message is					
TOLL FREE	not the intended recipient, you	not the intended recipient, you are hereby notified that any dissemination,					
Phone: 888.274.5158 Fax: 855.212.8	1, 0	distribution or copying of this communication is strictly prohibited. If you					
BCBSFL: 888.271.3183 Fax: 855.212.8 BCBSNJ: 888.214.1784 Fax: 855.212.8	I nave received this communicat	have received this communication in error, please return the original					
BCBSRI: 855.457.0759 Fax: 855.212.8	110 message to Prime Therapeutica	message to Prime Therapeutics via U.S. Mail. Thank you for your					
CHP: 855.457.0754 Fax: 855.212.8	cooperation.	cooperation.					