

Pharmacy Prior Authorization Form Fax completed form to: 877.974.4411 toll free, or 616.942.8206 ☐ Commercial (Traditional) ☐ Commercial (Individual/Optimized) This form applies to: Medicaid **Urgent** (life threatening) Non-**Urgent** (standard review) This request is: Urgent means the standard review time may seriously jeopardize the life or health of the patient or the patient's ability to regain maximum function. **Daraprim** (pyrimethamine) Member First Name: Last Name: DOB: _____ Gender: Primary Care Physician: Prov. Phone: Prov. Fax: Requesting Provider: Provider Address: Provider NPI: _____ Contact Name: Provider Signature: **Product Information** ☐ New request ☐ Continuation request Drug product: ☐ Daraprim 25mg Start date (or date of next dose): Date of last dose (if applicable): Dosing frequency: **Precertification Requirements** Before this drug is covered, the patient must meet one of the following requirements:

- 1. Treatment of toxoplasmosis
- 2. Secondary prevention of toxoplasmosis in patients with HIV
- 3. Prevention of pneumocystis pneumonia (PCP) in patients with HIV

If approved, the initial authorization will be for 6 weeks for toxoplasmosis and 3 months for pneumocystis. If approved for continuation, re-authorization may be required every 6 months for toxoplasmosis and every 3 months for pneumocystis.

For continuation when used for toxoplasmosis prophylaxis, patient must have met one of the following requirements:

- Patient remains symptomatic
- Patient is not receiving antiretroviral therapy
- Patient has a detectable HIV viral load
- Patient has maintained a CD4 count > 200 cells/microliter for less than six months

For continuation when used for pneumocystis prophylaxis, patient must have met one of the following requirements:

- CD4 count <200 cells/microliter
- Oropharyngeal candidiasis
- CD4 count percentage <14
- CD4 cell count between 200 and 250 cells/microliter IF frequent monitoring (eg, every three months) of CD4 cell counts is not possible



Note: Authorization for indications, dosing, or a route of administration not approved by the Food and Drug Administration (FDA) or recognized in CMS-accepted compendia (e.g. DrugDex, AHFS, U.S. Pharmacopeia, and also Clinical Pharmacology for oncology indications only) require supporting evidence for coverage. Please provide two published peer-reviewed literature articles supporting the appropriateness of the drug, or the route of administration to be used for the identified indication.

New request	
Priority Health Precertification Documentation	
Α.	What condition is this drug being requested for? Treatment of toxoplasmosis Secondary prevention of toxoplasmosis in patient with HIV Prevention of PCP (pneumocystis pneumonia) in patient with HIV Other – the patient's condition is: Rationale for use:
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Request to continue a previously authorized approval Priority Health Precertification Documentation	
Α.	What condition is this drug being requested for? Treatment of toxoplasmosis Secondary prevention of toxoplasmosis in patient with HIV Prevention of PCP (pneumocystis pneumonia) in patient with HIV Other – the patient's condition is:
<u>Co</u>	Rationale for use:
В.	Which of the following applies to the patient? Patient remains symptomatic Patient is not receiving antiretroviral therapy Patient has a detectable HIV viral load Patient has maintained a CD4 count > 200 cells/microliter for less than six months Other. Rationale for continued use:
Complete the following information for patients using Daraprim for pneumocystis prophylaxis: A. Which of the following applies to the patient?	
	 CD4 count <200 cells/microliter ☐ Oropharyngeal candidiasis ☐ CD4 count percentage <14 ☐ CD4 cell count between 200 and 250 cells/microliter IF frequent monitoring (eg, every three months) of CD4 cell counts is not possible ☐ Other. Rationale for continued use: