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AETNA BE	TTER HEALTH®						
Coverage Policy/Guideline							
Name: Daraprim (pyrimeth		namine)	Page:	1 of 2			
Effective Date: 6/26/2024			Last Review Date:	6/5/2024			
Applies to:	⊠Illinois	□Florida	□Michigan				
	□New Jersey	⊠Maryland	⊠Florida Kids				
	⊠Pennsylvania Kids	⊠Virginia					

Intent:

The intent of this policy/guideline is to provide information to the prescribing practitioner outlining the coverage criteria for Daraprim (pyrimethamine) under the patient's prescription drug benefit.

Description:

Daraprim (pyrimethamine) is indicated for the treatment of toxoplasmosis when used conjointly with a sulfonamide, since synergism exists with this combination.

Compendial Uses

Toxoplasmosis; Prophylaxis

Pneumocystis jirovecii pneumonia; Prophylaxis

Cystoisosporiasis; Treatment and secondary prophylaxis

Applicable Drug List:

Pyrimethamine

Policy/Guideline:

The requested drug will be covered with prior authorization when the following criteria are met:

 The requested drug is being prescribed for the treatment of congenital toxoplasmosis in a pediatric patient

OR

• The requested drug is being prescribed for the treatment of toxoplasmosis

OR

• The requested drug is being prescribed for secondary prophylaxis of toxoplasmosis

AND

 The patient has had a CD4 cell count of less than 200 cells/mm3 within the past 6 months

OR

 The patient has experienced an intolerance or has a contraindication to sulfamethoxazole/trimethoprim AND the requested drug is being prescribed for any of the following: A) primary prophylaxis of toxoplasmosis, B) *Pneumocystis jirovecii* pneumonia prophylaxis

AND

 The patient has had a CD4 cell count less than 200 cells/mm3 within the past 3 months

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 The patient has experienced an intolerance or has a contraindication to sulfamethoxazole/trimethoprim AND the requested drug is being prescribed for the treatment of cystoisosporiasis

OR

 The patient has experienced an intolerance or has a contraindication to sulfamethoxazole/trimethoprim AND the requested drug is being prescribed for secondary prophylaxis of cystoisosporiasis

AND

The patient has had a CD4 cell count less than 200 cells/mm3 within the past
6 months

Approval Duration and Quantity Restrictions:

- Treatment of congenital toxoplasmosis in a pediatric patient: 12 months
- Treatment of toxoplasmosis, primary prophylaxis of toxoplasmosis, or pneumocystis jirovecii pneumonia prophylaxis: 3 months
- Treatment of cystoisosporiasis, secondary prophylaxis of cystoisosporiasis, or secondary prophylaxis of toxoplasmosis: 6 months

References:

- 1. Daraprim [package insert]. New York, New York: Vyera Pharmaceuticals, LLC; August 2017.
- 2. Lexicomp Online, AHFS DI (Adult and Pediatric) Online. Waltham, MA: UpToDate, Inc.; 2023. https://online.lexi.com. Accessed December 11, 2023.
- 3. Micromedex (electronic version). Merative, Ann Arbor, Michigan, USA. Available at: https://www.micromedexsolutions.com/ (cited: 12/11/2023).
- 4. Panel on Guidelines for the Prevention and Treatment of Opportunistic Infections in Adults and Adolescents with HIV. Guidelines for the Prevention and Treatment of Opportunistic Infections in Adults and Adolescents with HIV. National Institutes of Health, Centers for Disease Control and Prevention, HIV Medicine Association, and Infectious Diseases Society of America. Available at https://clinicalinfo.hiv.gov/en/guidelines/adult-and-adolescent-opportunistic-infection. Accessed December 11,2023.
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- 6. Panel on Antiretroviral Guidelines for Adults and Adolescents. Guidelines for the Use of Antiretroviral Agents in Adults and Adolescents with HIV. Department of Health and Human Services. Available at https://clinicalinfo.hiv.gov/en/guidelines/adult-and-adolescent-arv. Accessed December 11, 2023.