

Drug Monograph

Drug/Drug **Daklinza[®] (daclatasvir) tablet/ Hepatitis C Anti-viral Agent**

Class:

Prepared for: MO HealthNet

Prepared by: Xerox Heritage, LLC

New Criteria

Revision of Existing Criteria

Executive Summary

Purpose:	The purpose of this monograph is to provide a review of new therapy to determine whether the reviewed drug should be made available on an open access basis to prescribers, require a clinical edit or require prior authorization for use.	
Dosage Forms:	Daklinza [®] is now available in a tablet containing 90 mg daclatasvir.	
Manufacturer:	Bristol-Myers Squibb Company, Princeton, NJ 08543	
Indications:	Daklinza [®] is indicated for use with sofosbuvir, with or without ribavirin, for the treatment of patients with chronic hepatitis C virus genotype 1 or genotype 3 infection.	
Costs:	\$757.50 per tablet of Daklinza [®] . <i>Maximum Allowable Cost</i>	
Summary of Findings:	This drug is being considered for inclusion in the state specific Preferred Drug List (PDL).	
Status Recommendation:	<input type="checkbox"/> Prior Authorization (PA) Required	<input type="checkbox"/> Open Access
	<input type="checkbox"/> Fiscal Edit	<input checked="" type="checkbox"/> PDL
Type of PA Criteria:	<input type="checkbox"/> Increased Risk of ADE	<input checked="" type="checkbox"/> Under Solicitation
	<input type="checkbox"/> Appropriate Indications	<input type="checkbox"/> No PA Required

Prepared By: Luke Boehmer Pharm.D.

Date: August 16, 2016