

Drug Monograph

Drug/Drug Class: **Diclegis™ (doxylamine succinate and pyridoxine hydrochloride) delayed release tablet/ antihistamine/B₆ analog**

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:	Diclegis™ is available as a delayed-release tablet containing 10 mg doxylamine succinate and 10 mg pyridoxine hydrochloride.	
Manufacturer:	Duchesnay USA, Inc., Bryn Mawr, PA, 19010,	
Indications:	Diclegis™ is a fixed dose combination drug product of doxylamine succinate, an antihistamine, and pyridoxine hydrochloride, a Vitamin B6 analog, indicated for the treatment of nausea and vomiting of pregnancy in women who do not respond to conservative management.	
Costs:	\$4.75 per delayed release tablet of Diclegis™ WholesaleAcquisitionCost	
Summary of Findings:	MO HealthNet Division recommends Prior Authorization status for this product.	
Status Recommendation:	<input checked="" type="checkbox"/> Prior Authorization (PA) Required	<input type="checkbox"/> Clinical Edit
	<input type="checkbox"/> Fiscal Edit	<input type="checkbox"/> PDL
Type of PA Criteria:	<input type="checkbox"/> Increased Risk of ADE	<input type="checkbox"/> Non-Preferred Agent
	<input checked="" type="checkbox"/> Appropriate Indications	<input type="checkbox"/> No PA Required

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