

Drug Monograph

Drug/Drug **Ibandronate sodium injection/ bisphosphonate**

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:	Ibandronate is available in a 3 mg/3 mL single dose vial containing 3 mg/3 mL ibandronate sodium for injection.	
Manufacturer:	Sun Pharmaceutical Industries Ltd. Acme Plaza, Andheri-Kurla Road Andheri (East), Mumbai - 400 059, India	
Indications:	Ibandronate sodium injection is a bisphosphonate indicated for the treatment of osteoporosis in postmenopausal women.	
Costs:	\$141.74 per vial of Ibandronate sodium <small>Maximum Allowable Cost</small>	
Summary of Findings:	This drug is being considered for inclusion in the state specific Preferred Drug List and is a non-preferred agent.	
Status Recommendation:	<input type="checkbox"/> Prior Authorization (PA) Required	<input type="checkbox"/> Clinical Edit
	<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"/> Non-Preferred Agent
	<input type="checkbox"/> Appropriate Indications	<input type="checkbox"/> No PA Required

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Date: May 16, 2014