

## Drug Monograph

Drug/Drug **Lanoxin™ (digoxin) tablet / cardiac glycoside**

Class:

Prepared for: MO HealthNet

Prepared by: Xerox Heritage, LLC

**New Criteria**

**Revision of Existing Criteria**

### Executive Summary

|                               |   |  |
|-------------------------------|---|--|
| <b>Purpose:</b>               | 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.           |  |
| <b>Dosage Forms:</b>          | Lanoxin™ is available in a new 187.5 mcg, unscored tablet containing 187.5 mg of digoxin.   |  |
| <b>Manufacturer:</b>          | DSM Pharmaceuticals, Inc.<br>Greenville, NC 27834   |  |
| <b>Indications:</b>           | Lanoxin™ is approved for the treatment of mild to moderate heart failure and to control the resting ventricular rate in chronic atrial fibrillation in adults as well as increasing myocardial contractility in pediatric patient with heart failure. |  |
| <b>Costs:</b>                 | \$2.40 per tablet of Lanoxin™ WholesaleAcquisitionCost  |  |
| <b>Summary of Findings:</b>   | MO HealthNet Division recommends Open Access status for this product.   |  |
| <b>Status Recommendation:</b> | <input type="checkbox"/> Prior Authorization (PA) Required  | <input checked="" type="checkbox"/> Open Access    |
|                               | <input type="checkbox"/> Fiscal Edit  | <input type="checkbox"/> PDL                       |
| <b>Type of PA Criteria:</b>   | <input type="checkbox"/> Increased Risk of ADE  | <input type="checkbox"/> Non-Preferred Agent       |
|                               | <input type="checkbox"/> Appropriate Indications  | <input checked="" type="checkbox"/> No PA Required |

Prepared By: Katie Wilbers, PharmD

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