# PREFERRED SPECIALTY MANAGEMENT POLICY

**POLICY:** Vesicular Monoamine Transporter Type 2 Inhibitors Preferred Specialty Management Policy

• Xenazine<sup>®</sup> (tetrabenazine tablets – Lundbeck, generic)

**REVIEW DATE:** 06/08/2022

#### **OVERVIEW**

Tetrabenazine, a reversible vesicular monoamine transporter type 2 (VMAT2) inhibitor, is indicated for the treatment of **chorea associated with Huntington's disease**.<sup>1</sup>

#### **Clinical Efficacy**

There are several published studies which have assessed the efficacy and safety of tetrabenazine for the treatment of other hyperkinetic movement disorders (e.g., tics in Tourette syndrome and tardive dyskinesia).<sup>2</sup>

#### **POLICY STATEMENT**

This Preferred Specialty Management program has been developed to encourage the use of the Preferred Product. For all medications (Preferred and Non-Preferred), the patient is required to meet the standard *Vesicular Monoamine Transporter Type 2 Inhibitors Prior Authorization Policy* criteria. The program also directs the patient to try the Preferred Product (generic tetrabenazine tablets). Requests for Non-Preferred Products will also be reviewed using the exception criteria (below). If the patient meets the standard *Prior Authorization Policy* criteria but has not tried a Preferred Product, approval for a Preferred Product will be authorized. All approvals are provided for 1 year.

**Documentation:** Documentation is required where noted in the criteria as **[documentation required]**. Documentation may include, but is not limited to, chart notes, prescription claims records, prescription receipts, and/or other information.

Automation: None

Preferred Product: Non-Preferred Product: Generic tetrabenazine tablets Xenazine

## **RECOMMENDED EXCEPTION CRITERIA**

Non-Preferred	Exception Criteria
Product	
Xenazine	<ol> <li>Patient meets the following criteria (A and B):         <ul> <li>A) Patient meets the standard Vesicular Monoamine Transporter Type 2 Inhibitor – Tetrabenazine Prior Authorization Policy criteria; AND</li> <li>B) Patient meets both of the following criteria (i and ii):                 <ul> <li>Patient tried generic tetrabenazine tablets; AND</li> <li>Bi Patient cannot continue to use the Preferred medication due to a formulation difference in the inactive ingredient(s) [e.g., difference in dyes, fillers, preservatives] between the Brand and the generic product which, per the prescribing physician, would result in a significant allergy or serious adverse reaction. [documentation required].</li> </ul> </li> <li>If the patient has met the standard Vesicular Monoamine Transporter Type 2 Inhibitor – Tetrabenazine Prior Authorization Policy criteria (1A), but has not met exception criteria (1B) above for brand Xenazine: approve generic</li> </ul></li></ol>
	<ul> <li>formulation difference in the inactive ingredient(s) [e.g., difference dyes, fillers, preservatives] between the Brand and the generic produ which, per the prescribing physician, would result in a significa allergy or serious adverse reaction. [documentation required].</li> <li>If the patient has met the standard <i>Vesicular Monoamine Transporter Type Inhibitor – Tetrabenazine Prior Authorization Policy</i> criteria (1A), but has <u>n</u></li> </ul>

### REFERENCES

1.

Xenazine<sup>®</sup> tablets [prescribing information]. Deerfield, IL: Lundbeck; September 2017. IBM Micromedex<sup>®</sup>. IBM Corporation. Available at: <u>www.micromedexsolutions.com</u>. Accessed on June 2, 2022. Search 2. terms: tetrabenazine.