Medimpact

STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

DEUTETRABENAZINE

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
DEUTETRABENAZINE	AUSTEDO,	44192		GPI-10	
	AUSTEDO XR			(6238003000)	

GUIDELINES FOR USE

- 1. Does the patient have a diagnosis of chorea (involuntary movements) associated with Huntington's disease and meet **ALL** of the following criteria?
 - The patient is 18 years of age or older
 - Therapy is prescribed by or in consultation with a neurologist or movement disorder specialist

If yes, approve for 12 months by GPID or GPI-14 for ALL of the following:

- 6-12-24mg XR titration kit: #42 per 28 days for 1 fill.
- 6mg: #2 per day.
- 9mg: #4 per day.
- 12mg: #4 per day.
- 6mq XR: #7 per day.
- 12mg XR: #3 per day.
- 24mg XR: #2 per day.

If no, continue to #2.

- 2. Does the patient have a diagnosis of moderate to severe tardive dyskinesia (TD) and meet ALL of the following criteria?
 - The patient is 18 years of age or older
 - The patient's TD has been present for at least 3 months
 - Therapy is prescribed by or in consultation with a neurologist, movement disorder specialist, or psychiatrist
 - The patient has a prior history of using antipsychotic medications (e.g., aripiprazole, haloperidol, ziprasidone) or metoclopramide for at least 3 months (or at least 1 month if patient is 60 years of age or older) as documented in the prescription claims history

If yes, approve for 12 months by GPID or GPI-14 for ALL of the following:

- 6-12-24mg XR titration kit: #42 per 28 days for 1 fill.
- 6mg: #2 per day.
- 9mg: #4 per day.
- 12mg: #4 per day.
- 6mg XR: #7 per day.
- 12mg XR: #3 per day.
- 24mg XR: #2 per day.

If no, do not approve.

DENIAL TEXT: See the denial text at the end of the guideline.

CONTINUED ON NEXT PAGE

Copyright © 2023 MedImpact Healthcare Systems, Inc. All rights reserved. This document is proprietary to MedImpact. MedImpact maintains the sole and exclusive ownership, right, title, and interest in and to this document.

Medimpact

STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

DEUTETRABENAZINE

GUIDELINES FOR USE (CONTINUED)

DENIAL TEXT: *Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

Our guideline named **DEUTETRABENAZINE (Austedo)** requires the following rule(s) be met for approval:

- A. You have ONE of the following diagnoses:
 - 1. Chorea (involuntary muscle movements) associated with Huntington's disease
 - 2. Moderate to severe tardive dyskinesia (uncontrolled body movements)
- B. You are 18 years of age or older
- C. If you have chorea associated with Huntington's disease, approval also requires:
 - 1. Therapy is prescribed by or in consultation with a neurologist (type of brain doctor) or movement disorder specialist
- D. If you have moderate to severe tardive dyskinesia, approval also requires:
 - 1. Moderate to severe tardive dyskinesia (uncontrolled body movements) has been present for at least 3 months
 - 2. Therapy is prescribed by or in consultation with a neurologist (type of brain doctor), movement disorder specialist, or psychiatrist (type of mental health doctor)
 - 3. You have a prior history of using antipsychotic medications (such as aripiprazole, haloperidol, ziprasidone) or metoclopramide for at least 3 months (or at least 1 month if you are 60 years of age or older) as documented in your prescription claims history

Your doctor told us [INSERT PT SPECIFIC INFO PROVIDED]. We do not have information showing you [INSERT UNMET CRITERIA]. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

RATIONALE

For further information, please refer to the Prescribing Information and/or Drug Monograph for Austedo, Austedo XR.

REFERENCES

 Austedo, Austedo XR [Prescribing Information]. Parsippany, NJ: Teva Neuroscience, Inc.; February 2023.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A Commercial Effective: 08/21/23 Created: 04/17 Client Approval: 07/23

P&T Approval: 01/22

Copyright © 2023 MedImpact Healthcare Systems, Inc. All rights reserved. This document is proprietary to MedImpact. MedImpact maintains the sole and exclusive ownership, right, title, and interest in and to this document.