

AUSTEDO (deutetrabenazine)

Pre - PA Allowance

None

Prior-Approval Requirements

Age 18 years of age or older

Diagnoses

Patient must have **ONE** of the following:

1. Moderate to severe tardive dyskinesia

AND ALL of the following:

- a. Inadequate treatment response, intolerance, or contraindication to **ONE** of the following:
 - i. Benzodiazepine
 - ii. Second generation antipsychotic (e.g., Seroquel, clozapine)
 - iii. Xenazine
- b. Documented baseline evaluation of the condition using **ONE** of the following scoring tools:
 - i. Abnormal Involuntary Movement Scale (AIMS)
 - ii. Extrapyramidal Symptom Rating Scale (ESRS)
- c. Prescriber has reduced the dosage or discontinued all causative medications including antipsychotic medication and metoclopramide (Reglan)
- d. Patient has a functional impairment that justifies treatment with Austedo
- 2. Chorea associated with Huntington's disease

AND NONE of the following for ALL indications:

- 1. Actively suicidal
- 2. Untreated or inadequately treated depression
- 3. Concomitant use of a MAOI (monoamine oxidase inhibitor) (must be >14 days post discontinuing therapy)
- 4. Concomitant use of reserpine (must be >20 days post discontinuing therapy)
- 5. Hepatic impairment
- 6. Dual therapy with other vesicular monoamine transporter 2 (VMAT2) inhibitors



AUSTEDO (deutetrabenazine)

Prior - Approval Limits

Quantity48mg per dayDuration12 months

Prior – Approval Renewal Requirements

Age: 18 years of age or older

Diagnoses

Patient must have ONE of the following:

1. Tardive dyskinesia

AND the following:

- a. Documented improvement using **ONE** of the following scores:
 - i. Abnormal Involuntary Movement Scale (AIMS)
 - ii. Extrapyramidal Symptom Rating Scale (ESRS)
- 2. Chorea associated with Huntington's disease

AND NONE of the following for ALL indications:

- 1. Actively suicidal
- 2. Untreated or inadequately treated depression
- 3. Concomitant use of a MAOI (monoamine oxidase inhibitor) (must be >14 days post discontinuing therapy)
- Concomitant use of reserpine (must be >20 days post discontinuing therapy)
- 5. Hepatic impairment
- 6. Dual therapy with other vesicular monoamine transporter 2 (VMAT2) inhibitors

Prior - Approval Renewal Limits

Same as above