

VMAT2 INHIBITORS PRIOR AUTHORIZATION FORM (form effective 2/1/2022)

Prior authorization guidelines for **VMAT2 Inhibitors** and **Quantity Limits/Daily Dose Limits**
 are available on Geisinger Health Plan's website at
<https://healthplan.geisinger.org/pharmacy/pharmacy.aspx?strip=true&style=OneGeisinger>

<input type="checkbox"/> New request <input type="checkbox"/> Renewal request		Total # of pages: _____		Prescriber name:	
Name of office contact:				Specialty:	
Contact's phone number:				NPI:	State license #:
LTC facility contact/phone:				Street address:	
Beneficiary name:				Suite #:	City/state/zip:
Beneficiary ID#:		DOB:		Phone:	Fax:
Beneficiary address:				Beneficiary phone number:	

CLINICAL INFORMATION

Drug requested: <input type="checkbox"/> Austedo tablet <input type="checkbox"/> tetrabenazine tablet <input type="checkbox"/> Ingrezza capsule <input type="checkbox"/> Xenazine tablet (<i>submit documentation showing why generic tetrabenazine cannot be used</i>) <input type="checkbox"/> Ingrezza initiation pack <input type="checkbox"/> _____		
Strength:	Quantity:	Refills:
Dose/directions:		
Diagnosis (<u>submit documentation</u>):		DX codes (<u>required</u>):
Is the requested medication being prescribed by or in consultation with a specialist (ie., neurologist or psychiatrist)?		<input type="checkbox"/> Yes <i>Submit documentation of consultation</i> <input type="checkbox"/> No <i>if applicable.</i>

Check all of the following that apply to the beneficiary and this request and SUBMIT DOCUMENTATION for each item.

☐ **For AUSTEDO (DEUTETRABENAZINE):**

- ☐ Has a contraindication to Austedo (*check all that apply*):
- ☐ Actively suicidal
 - ☐ Taken an MAO inhibitor in the past 14 days (eg, phenelzine, rasagiline, selegiline, tranylcypromine)
 - ☐ Hepatic impairment
 - ☐ Taken reserpine in the past 20 days
 - ☐ Taking Xenazine or Ingrezza
 - ☐ Depression that is untreated or inadequately treated
- ☐ Is known to be a poor CYP2D6 metabolizer
- ☐ Austedo dose is adjusted accordingly based on dosing recommendations in the package labeling
- ☐ Will be taking a strong CYP2D6 inhibitor while taking Austedo (e.g., bupropion, fluoxetine, paroxetine, quinidine)
- ☐ Austedo dose is adjusted accordingly based on dosing recommendations in the package labeling

☐ **For INGREZZA (VALBENAZINE):**

- ☐ Is taking a strong CYP3A4 inhibitor (eg, some azole antifungals, nefazodone, some protease inhibitors)
- ☐ Ingrezza dose is adjusted accordingly based on dosing recommendations in the package labeling

- ☐ Is taking a strong CYP2D6 inhibitor (eg, bupropion, fluoxetine, paroxetine, quinidine)
- ☐ Ingrezza dose is adjusted accordingly based on dosing recommendations in the package labeling
- ☐ Has moderate or severe hepatic impairment (Child-Pugh score 7 to 15)
- ☐ Ingrezza dose is adjusted accordingly based on dosing recommendations in the package labeling
- ☐ Is taking a strong CYP3A4 inducer (eg, carbamazepine, phenytoin, rifampin, St. John's Wort) (*concomitant use not recommended per package labeling*)

☐ **For TETRABENAZINE / XENAZINE:**

- ☐ Has a contraindication to tetrabenazine / Xenazine (*check all that apply*):
- ☐ Actively suicidal
 - ☐ Taken an MAO inhibitor in the past 14 days (eg, phenelzine, rasagiline, selegiline, tranylcypromine)
 - ☐ Hepatic impairment
 - ☐ Taken reserpine in the past 20 days
 - ☐ Taking Austedo or Ingrezza
 - ☐ Depression that is untreated or inadequately treated
- ☐ Will be taking a strong CYP2D6 inhibitor while taking tetrabenazine (eg, bupropion, fluoxetine, paroxetine, quinidine)
- ☐ Tetrabenazine dose is adjusted accordingly based on dosing recommendations in the package labeling
- ☐ Is prescribed a tetrabenazine dose that exceeds 50 mg per day
- ☐ Has documentation of therapeutic failure of tetrabenazine at a dose of ≤ 50 mg/day
- ☐ Has documentation of CYP450 2D6 genotyping showing intermediate or extensive metabolism

INITIAL requests

Check all of the following that apply to the beneficiary and this request and SUBMIT DOCUMENTATION for each item.

- ☐ Had a mental health evaluation
- ☐ Has a history of suicide attempt, bipolar disorder, or major depressive disorder
 - ☐ Was evaluated in the past 6 months and treated by a psychiatrist
 - ☐ Was determined to be a candidate for treatment with the requested medication based on the mental health evaluation
- ☐ **For treatment of TARDIVE DYSKINESIA:**
- ☐ Has no other causes of involuntary movement
 - ☐ Has documentation of tardive dyskinesia severity using a validated scale or assessment of impact on daily function
 - ☐ A decrease in dose of dopamine receptor blocking agents is not appropriate

RENEWAL requests

Check all of the following that apply to the beneficiary and this request and SUBMIT DOCUMENTATION for each item.

- ☐ Was reevaluated for new onset or worsening symptoms of depression
- ☐ If applicable, was or is being treated for symptoms of depression
- ☐ Was determined to remain a candidate for treatment with the requested medication based on the mental health evaluation
- ☐ For treatment of CHOREA:
- ☐ Experienced clinical benefit from the requested medication based on the prescriber's clinical judgement
- ☐ For treatment of TARDIVE DYSKINESIA:
- ☐ Experienced an improvement in tardive dyskinesia severity documented by a validated scale
- ☐ Experienced an improvement in daily functioning

Please submit to PromptPA <https://ghp.promptpa.com> OR fax to Geisinger Health Plan at 570-271-5610 the completed form with required clinical documentation.

Prescriber Signature:

Date:

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