

HEALTHY U MEDICAID

PRIOR AUTHORIZATION REQUEST FORM

VMAT-2 INHIBITORS

Austedo®, Ingrezza®, tetrabenazine

For authorization, please answer each question and fax this form PLUS chart notes back to the Healthy U Prior Authorization Department at 888-509-8142.

Failure to submit clinical documentation to support this request will result in a dismissal of the request.

If you have prior authorization questions, please call for Pharmacy Customer Service for assistance at 855-856-5694

Disclaimer: Prior authorization request forms are subject to change in accordance with Federal and State notice requirements.

Date:	Member Name:	ID#:
DOB:	Gender:	Physician:
Office Phone:	Office Fax:	Office Contact:
Height/Weight:		

Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria.

Product being requested: Austedo® (deutetrabenazine), Ingrezza® (valbenazine), tetrabenazine

Dosing/Frequency: _____

If the request is for reauthorization, proceed to reauthorization section

Questions	Yes	No	Comments/Notes
TARDIVE DYSKINESIA			
1. Is the request made by, or in consultation with, a psychiatrist or neurologist?	<input type="checkbox"/>	<input type="checkbox"/>	
2. Does documentation contain an Abnormal Involuntary Movement Scale (AIMS)?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
3. Does documentation contain a Clinical Global Impression of Severity (CGI-S)?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
4. Is the member currently taking reserpine or a MAO-I?	<input type="checkbox"/>	<input type="checkbox"/>	
5. For Ingrezza®, has the member had a 3-month trial and failure of a benzodiazepine and tetrabenazine?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
6. For Austedo®, has the member had a 3-month trial and failure of Ingrezza®?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
CHOREA ASSOCIATED WITH HUNTINGTON'S DISEASE			
1. Is the request made by, or in consultation with, a neurologist?	<input type="checkbox"/>	<input type="checkbox"/>	
2. Does documentation include a diagnosis made by characteristic motor examination features and genetic testing?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
3. Is the member ambulatory?	<input type="checkbox"/>	<input type="checkbox"/>	
4. Has the member tried and failed at least two of the following: amantadine, an antipsychotic, riluzole, nabilone?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation

5. For Austedo®, has the member had a 3-month trial and failure of tetrabenazine?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
TOURETTE SYNDROME			
1. Have non-pharmacologic therapies been tried and found to be inadequate to meet treatment goals?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
2. Has the member tried and failed at least 3 of the following medications: guanfacine, clonidine, topiramate, botulinum toxin, fluphenazine, haloperidol, risperidone, pimozide, aripiprazole?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
REAUTHORIZATION			
1. Does clinical documentation show a continued medical need as well as medication efficacy defined as objective progress towards treatment plan goals?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
2. If the request for reauthorization of therapy for tardive dyskinesia, does clinical documentation show a continued medical need as well as medication efficacy defined by the AIMS score has decreased by at least 2 points from based line and the CGI-S score is ≤ 2 ?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
3. If the request is for reauthorization of therapy for Huntington's disease, does updated documentation show disease stabilization and functional improvement of symptoms?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
What medications and/or treatment modalities have been tried in the past for this condition? Please document name of treatment, reason for failure, treatment dates, etc.			
Additional information:			
Physician's Signature:			

**** Failure to submit clinical documentation to support this request will result in a dismissal of the request. ****

Policy: PHARM-HU-077
 Origination Date: 01/01/2022
 Reviewed/Revised Date: 05/18/2022
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