

Please complete this **entire** form and fax it to: **866-940-7328**. If you have questions, please call **800-310-6826**.
This form may contain multiple pages. Please complete all pages to avoid a delay in our decision. Allow at least 24 hours for review.

Apple Health Preferred Drug list: <https://www.hca.wa.gov/assets/billers-and-providers/apple-health-preferred-drug-list.xlsx>

Section A – Member Information

First Name:	Last Name:	Member ID:
Address:		
City:	State:	ZIP Code:
Phone:	DOB:	Allergies:
Primary Insurance Information (if any):		
Is the requested medication: <input type="checkbox"/> New or <input type="checkbox"/> Continuation of Therapy? If continuation, list start date: _____		
Is this patient currently hospitalized? <input type="checkbox"/> Yes <input type="checkbox"/> No If recently discharged, list discharge date: _____		

Section B - Provider Information

First Name:	Last Name: _____ M.D./D.O.		
Address:	City:	State:	ZIP code:
Phone:	Fax:	NPI #:	Specialty:
Office Contact Name / Fax attention to:			

Section C - Medical Information

Medication:	Strength:
Directions for use:	Quantity:
Diagnosis (Please be specific & provide as much information as possible):	ICD-10 CODE:
Is this member pregnant? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, what is this member's due date? _____	

Section D – Previous Medication Trials

Medication Name	Strength	Directions	Dates of Therapy	Reason for failure / discontinuation

**Section E – Additional information and Explanation of why preferred medications would not meet the patient's needs:
Please refer to the patient's PDL for a list of preferred alternatives**

1. Is this request for a continuation of therapy? Yes No
 If yes, does patient have clinical documentation demonstrating disease stability or a positive clinical response (e.g, reduction in involuntary movements, decrease in total maximal chorea score, or improvement in AIMS or CGI-TD score)?
 Yes No
2. Indicate patient's diagnosis:
 Chorea associated with Huntington's disease
 Tardive dyskinesia
 Other. Specify: _____
3. Is this being prescribed by or in consultation with a psychiatrist or neurologist? Yes No
4. Has a baseline assessment been completed using any of the following? Check all that apply:
 The Unified Huntington's Disease Rating Scale (UHDRS). Specify section completed:
 Motor
 Cognitive
 Behavioral
 Functional Assessment
 Independence Scale
 Total Functional Capacity
 Abnormal Involuntary Movement Scale (AIMS)
 Clinical Global Impression of Change – Tardive Dyskinesia (CGI-TD)
5. Will this be used in combination with a monoamine oxidase inhibitor (MAOI) [e.g. isocarboxazid, phenelzine, tranylcypromine, reserpine] or another vesicular monoamine transporter 2 (VMAT2) inhibitor [e.g. tetrabenazine]? Yes No
6. Has patient had treatment (minimum of 12 weeks) with deutetrabenazine or deutetrabenazine ER that has been ineffective, not tolerated, or is treatment contraindicated? Yes No
7. What alternative treatments has patient tried?
 What was the outcome of the trial?

If patient has tardive dyskinesia:

8. Does patient continue to experience persistent TD after trying one of the following unless contraindicated, not tolerated, or put patient's psychiatric stability at risk?
 Switching from a 1st generation to a 2nd generation antipsychotic
 Patient has tried two 2nd generation antipsychotics
 Patient has a history of discontinuation or dose modification of the offending medication with continued symptoms

CHART NOTES ARE REQUIRED WITH THIS REQUEST

Prescriber signature

Prescriber specialty

Date