

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.

| Member Information | | | Prescriber Information | | |
|---|--------|------------|------------------------|-----------|------------|
| Member Name: | | | Provider Name: | | |
| Member ID: | | | NPI #: | | Specialty: |
| Date Of Birth: | | | Office Phone: | | |
| Street Address: | | | Office Fax: | | |
| City: | State: | ZIP Code: | Office Street Address: | | |
| Phone: | | Allergies: | City: | State: | ZIP Code: |
| 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: _____ Is this member pregnant? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, what is this member's due date? _____ | | | | | |
| Medication Information | | | | | |
| Medication: | | | | Strength: | |
| Directions for use: | | | | Quantity: | |
| Medication Administered: <input type="checkbox"/> Self-Administered <input type="checkbox"/> Physician's Office <input type="checkbox"/> Other: _____ | | | | | |
| Clinical Information | | | | | |
| What is the patient's diagnosis for the medication being requested? _____ _____ | | | | | |
| ICD-10 Code(s): _____ | | | | | |
| Are there any supporting laboratory or test results related to the patient's diagnosis? <i>(Please specify or provide documentation)</i> | | | | | |
| Previous Medication Trials / Contraindications | | | | | |
| <u>Please refer to the patient's PDL at www.uhcprovider.com for a list of preferred alternatives</u> | | | | | |
| What medication(s) does the patient have a history of <u>failure</u> to? <i>(Please specify ALL medication(s)/strengths tried, directions, length of trial, and reason for discontinuation of each medication)</i> | | | | | |
| What medication(s) does the patient have a <u>contraindication or intolerance</u> to? <i>(Please specify ALL medication(s) with the associated contraindication to or specific issues resulting in intolerance to each medication)</i> | | | | | |
| Additional information that may be important for this review | | | | | |
| | | | | | |

| | | |
|--------------------|-------------------|-------------|
| Member First name: | Member Last name: | Member DOB: |
|--------------------|-------------------|-------------|

Clinical and Drug Specific Information

ALL REQUESTS

| | |
|---|--|
| <input type="checkbox"/> Yes <input type="checkbox"/> No | Does the patient have a diagnosis of tardive dyskinesia? |
| <input type="checkbox"/> Yes <input type="checkbox"/> No | Is Ingrezza prescribed by or in consultation with a neurologist or a psychiatrist? |
| <input type="checkbox"/> Yes <input type="checkbox"/> No | Does the patient have a contraindication to Ingrezza? |
| <input type="checkbox"/> Yes <input type="checkbox"/> No | Does the patient have a history of prior suicide attempt, bipolar disorder, or major depressive disorder? |
| <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Not applicable | If yes to the above question, was the patient evaluated within the previous 6 months and treated by a psychiatrist? |
| <input type="checkbox"/> Yes <input type="checkbox"/> No | Has the patient had a mental health evaluation performed? |
| <input type="checkbox"/> Yes <input type="checkbox"/> No | Does the patient have any of the following? <i>(If yes, check which applies)</i> <ul style="list-style-type: none"> <input type="checkbox"/> Was assessed for and determined to have no other causes of involuntary movement <input type="checkbox"/> Was evaluated for appropriateness of dose decrease of dopamine receptor blocking agents <input type="checkbox"/> Has documentation of tardive dyskinesia severity using a validated scale or assessment of impact on daily function |

CONTINUATION OF THERAPY

| | |
|--|---|
| <input type="checkbox"/> Yes <input type="checkbox"/> No | Has the patient experienced an improvement in tardive dyskinesia severity documented by a validated scale or improvement in daily function? |
| <input type="checkbox"/> Yes <input type="checkbox"/> No | Was the patient re-evaluated and treated for new onset or worsening symptoms of depression and determined to continue to be a candidate for treatment with Ingrezza? |

Provider Signature: _____ **Date:** _____

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