

## NC Pharmacy Prior Approval Request for Antiparkinson's Agents-Inbrija and Ongentys

### Beneficiary Information

1. Beneficiary Last Name: \_\_\_\_\_ 2. First Name: \_\_\_\_\_  
3. Beneficiary ID #: \_\_\_\_\_ 4. Beneficiary Date of Birth: \_\_\_\_\_ 5. Beneficiary Gender: \_\_\_\_\_

### Prescriber Information

6. Prescribing Provider NPI #: \_\_\_\_\_ Provider Fax #: \_\_\_\_\_  
7. Requester Contact Information - Name: \_\_\_\_\_ Phone #: \_\_\_\_\_ Ext. \_\_\_\_\_

### Drug Information

8. Drug Name: \_\_\_\_\_ 9. Strength: \_\_\_\_\_ 10. Quantity Per 30 Days: \_\_\_\_\_  
11. Length of Therapy (in days):  up to 30 Days  60 Days  90 Days  120 Days  180 Days  365 Days

### Clinical Information

#### **Inbrija - initial authorization requests **\*\*Initial requests can be approved for up to 6 months\*\***:**

1. Is the beneficiary age 18 or older?  **Yes**  **No**
2. Does the beneficiary have a diagnosis of Parkinson's Disease and is experiencing "off" episodes?  **Yes**  **No**
3. Will the beneficiary be concurrently receiving optimized carbidopa/levodopa therapy?  **Yes**  **No**
4. Is the beneficiary currently taking a nonselective monoamine (MAO) inhibitor or has the beneficiary taken a MAO inhibitor within the last two weeks?  **Yes**  **No**
5. Does the beneficiary have asthma, COPD or other chronic lung disease?  **Yes**  **No**

#### **Inbrija - reauthorization requests (please answer questions 1-6) **\*\*Reauthorization requests can be approved for up to 12 months\*\***:**

6. Has documentation been submitted that indicates the beneficiary has had an improvement in their symptoms from baseline?  **Yes**  **No**

#### **Ongentys - initial authorization requests **\*\*Initial requests can be approved for up to 6 months\*\***:**

7. Is the beneficiary age 18 years of age or older?  **Yes**  **No**
8. Does the beneficiary have a diagnosis of Parkinson's Disease and is experiencing "off" episodes for at least 1.5 hours/day on average?  **Yes**  **No**
9. Does the beneficiary have no contraindications including ESRD (creatinine clearance <15 ml/min/1.73m<sup>2</sup>)?  
 **Yes**  **No**
10. Does the beneficiary have no contraindications including severe hepatic impairment (Child-Pugh C)?  
 **Yes**  **No**
11. Is the beneficiary currently taking a nonselective monoamine oxidase-B (MAO-B) inhibitor?  **Yes**  **No**
12. Will the beneficiary be concurrently receiving optimized carbidopa/levodopa therapy?  **Yes**  **No**
13. Has the beneficiary had an adequate trial and subsequent failure of at least 2 preferred adjunctive therapies (e.g., dopamine agonists, MAO-B inhibitors, catechol-O-methyltransferase [COMT] inhibitors) to control "off" symptoms?  **Yes**  **No**

#### **Ongentys - reauthorization requests (please answer questions 7-15) **\*\*Reauthorization requests can be approved for up to 12 months\*\***:**

14. Has documentation been submitted that indicates the beneficiary has had clinically meaningful response to treatment (e.g., beneficiary shows a reduction in time of "off" episodes)?  **Yes**  **No**
15. Has the beneficiary experienced toxicity or treatment related adverse event from the drug (e.g., dyskinesias, hallucinations/psychotic behavior, impulse control/compulsive behaviors)?  **Yes**  **No**

Signature of Prescriber: \_\_\_\_\_ Date: \_\_\_\_\_

**(Prescriber Signature Mandatory)**

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.