

# OPTIMA HEALTH PLAN

## PHARMACY/MEDICAL PRIOR AUTHORIZATION/STEP-EDIT REQUEST\*

Directions: The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; fax to 1-844-202-5034. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct.

**Drug Requested:**                    **Immune Globulin Intravenous (IVIG) -**  
**(Multifocal Motor Neuropathy-MMN)**

**DRUG INFORMATION:** Complete information below. If incomplete, authorization process will be delayed.

**Circle J Code that applies:**        **J1459 / J1556 / J1561 / J1566 / J1568 / J1569 / J1572**

**Drug Form/Strength/Month:** \_\_\_\_\_

**Dosing Schedule:** \_\_\_\_\_                    **Length of Therapy:** \_\_\_\_\_

**Diagnosis:** \_\_\_\_\_                    **ICD Code:** \_\_\_\_\_

**\*Medical notes must be submitted to support each line checked on this request.\***

**CLINICAL DIAGNOSIS/CRITERIA:** Check one of the applicable diagnoses below. Boxes **MUST** be checked to qualify. If incomplete, authorization process will be delayed.

- Multifocal Motor Neuropathy (MMN): initial trial 4 weeks: (Check one of the following):**
  - Asymmetric weakness that affects distal muscles  Yes  No
  - Does the patient have upper motor neuron signs?  Yes  No
  - Does the nerve conduction study confirm a demyelinating neuropathy is present (conduction block, slowing, or abnormal temporal dispersion in at least one nerve)?  Yes  No
- OR**
- History and exam do not suggest upper motor neuron disease (*no bulbar weakness, no upper motor neuron signs*)  Yes  No
- Labs show that GM-1 antibody titers are elevated  Yes  No
- OR**
- Electrodiagnostic testing clinical presentation suggests MMN but the diagnosis remains uncertain  Yes  No
- Continued use of Ig after initial trial for MMN when the following criteria are met:**
  - Progress notes document an improvement in strength and function within three weeks of the start of the infusion period  Yes  No
  - Continue need if during annual basis the dose was titrated or change in interval of therapy result in worsening of symptoms

(signature on next page)

**Medication being provided by (check applicable box below):**

Physician's office

**OR**

Specialty Pharmacy:

Briova SpecialtyRx

**\*\*Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.\*\***

**\*Previous therapies will be verified through pharmacy paid claims or submitted chart notes.\***

Patient Name: \_\_\_\_\_

Member Optima #: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

Prescriber Name: \_\_\_\_\_

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

Office Contact Name: \_\_\_\_\_

Phone Number: \_\_\_\_\_ Fax Number: \_\_\_\_\_

DEA OR NPI #: \_\_\_\_\_

\*Approved by Pharmacy and Therapeutics Committee: 4/15/2013

REVISED/UPDATED: 6/30/2013; 8/19/2014; 10/31/2014; 5/23/2015; 12/30/2015; 1/29/2016; 8/18/2016; 9/22/2016; 11/29/2016; 12/13/2016; 9/15/2017; 10/6/2017