# SPECIALTY GUIDELINE MANAGEMENT

# COPAXONE (glatiramer acetate) GLATOPA (glatiramer acetate) glatiramer acetate

### **POLICY**

## I. INDICATIONS

The indications below including FDA-approved indications and compendial uses are considered a covered benefit provided that all the approval criteria are met and the member has no exclusions to the prescribed therapy.

# **FDA-Approved Indication**

Indicated for the treatment of patients with relapsing forms of multiple sclerosis (MS), to include clinically isolated syndrome, relapsing-remitting disease, and active secondary progressive disease, in adults.

All other indications are considered experimental/investigational and not medically necessary.

### II. PRESCRIBER SPECIALTIES

This medication must be prescribed by or in consultation with a neurologist.

# III. CRITERIA FOR INITIAL APPROVAL

### A. Relapsing forms of multiple sclerosis

Authorization of 12 months may be granted to members who have been diagnosed with a relapsing form of multiple sclerosis (including relapsing-remitting and secondary progressive disease for those who continue to experience relapse).

# B. Clinically isolated syndrome

Authorization of 12 months may be granted to members for treatment of clinically isolated syndrome of multiple sclerosis.

# IV. CONTINUATION OF THERAPY

For all indications: Authorization of 12 months may be granted for members who are experiencing disease stability or improvement while receiving Copaxone, Glatopa, or glatiramer acetate.

# V. OTHER

Members will not use Copaxone, Glatopa, or glatiramer acetate concomitantly with other disease modifying multiple sclerosis agents (Note: Ampyra and Nuedexta are not disease modifying).

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## VI. REFERENCES

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- 2. Glatopa [package insert]. Princeton, NJ: Sandoz Inc.; December 2023.
- 3. Glatiramer acetate 20mg/mL [package insert]. Morgantown, WV: Mylan Pharmaceuticals Inc.; January 2024.
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