	TTER HEALTH® Policy/Guideline	<b>*ae</b>	etna™	
Name:	Aubagio		Page:	1 of 2
Effective Date: 11/1/2024			Last Review Date:	10/2024
Applies	□Illinois	□Florida ⊠Manuland	⊠Florida Kids	
to:	⊠New Jersey ⊠Pennsylvania Kids	⊠Maryland □Virginia	□Michigan □Texas	

## Intent:

The intent of this policy/guideline is to provide information to the prescribing practitioner outlining the coverage criteria for Aubagio (teriflunomide) under the patient's prescription drug benefit.

# **Description:**

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**

Aubagio is 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.

# **Applicable Drug List:**

Teriflunomide

### **Policy/Guideline:**

### **Prescriber Specialty:**

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

# 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 the treatment of clinically isolated syndrome of multiple sclerosis.

### **Continuation of Therapy:**

	ETTER HEALTH® Policy/Guideline	♥aetna™		
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For all indications: Authorization of 12 months may be granted to members who are experiencing disease stability or improvement while receiving Aubagio.

## **Other Criteria:**

- A. Members will not use Aubagio concomitantly with other disease modifying multiple sclerosis agents (Note: Ampyra and Nuedexta are not disease modifying).
- B. Authorization may be granted for pediatric members less than 18 years of age when benefits outweigh risks.

## Approval Duration and Quantity Restrictions:

Approval: 12 months

Quantity Level Limit: 30 tablets per 30 days

#### **References:**

- 1. Aubagio [package insert]. Cambridge, MA: Genzyme Corporation; June 2024.
- 2. Teriflunomide [package insert]. East Windsor, NJ: Aurobindo Pharma USA, Inc.; February 2024.