



STANDARD COMMERCIAL DRUG FORMULARY
PRIOR AUTHORIZATION GUIDELINES

INTERFERON FOR MS - EXTAVIA

| Generic | Brand | HICL | GCN | Medi-Span | Exception/Other |
|--------------------|---------|-------|-----|---------------------|-----------------|
| INTERFERON BETA-1B | EXTAVIA | 08537 | | GPI-10 (6240306050) | BRAND = EXTAVIA |

GUIDELINES FOR USE

1. Does the patient have a diagnosis of a relapsing form of multiple sclerosis (MS), to include clinically isolated syndrome, relapsing remitting disease, and active secondary progressive disease and meet **ALL** of the following criteria?
 - The patient is 18 years of age or older
 - The patient had a trial of or contraindication to any TWO of the following preferred agents for MS: Avonex, Copaxone/Glatiramer/Glatopa, Gilenya, Plegridy, Rebif, Betaseron, dimethyl fumarate, Mavenclad, Mayzent, Vumerity, Aubagio, Kesimpta (Please note: other multiple sclerosis agents may also require prior authorization)

If yes, **approve for 12 months for all NDCs or GPI-14 of Extavia for #14 vials or kits per 28 days.**

If no, do not approve.

DENIAL TEXT: *Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

Our guideline named **INTERFERON FOR MS - EXTAVIA** requires the following rule(s) be met for approval:

- A. You have a relapsing form of multiple sclerosis (MS: a type of nerve disorder), to include clinically isolated syndrome (disease occurs once), relapsing-remitting disease (symptoms go away and return), and active secondary progressive disease (advanced disease)
- B. You are 18 years of age or older
- C. You had a trial of or contraindication (harmful for) to any TWO of the following preferred medications: Avonex, Copaxone/Glatiramer/Glatopa, Gilenya, Plegridy, Rebif, Betaseron, dimethyl fumarate, Mavenclad, Mayzent, Vumerity, Aubagio, Kesimpta (Please note: other multiple sclerosis medications may also require prior authorization)

Your doctor told us **[INSERT PT SPECIFIC INFO PROVIDED]**. We do not have information showing you **[INSERT UNMET CRITERIA]**. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

CONTINUED ON NEXT PAGE



**STANDARD COMMERCIAL DRUG FORMULARY
PRIOR AUTHORIZATION GUIDELINES**

INTERFERON FOR MS - EXTAVIA

RATIONALE

For further information, please refer to the Prescribing Information and/or Drug Monograph for Extavia.

REFERENCES

- Extavia [Prescribing Information]. East Hanover, NJ: EMD Novartis; August 2019.

| | | |
|---------|------------|-----|
| Library | Commercial | NSA |
| Yes | Yes | No |

Part D Effective: N/A

Commercial Effective: 11/01/22

Created: 10/22

Client Approval: 10/22

P&T Approval: 01/20