

## AUBAGIO Prior Authorization Form

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- Standard Request
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- Expedited Request

If you or your prescriber believe that waiting for a standard decision could seriously harm your life, health, or ability to regain maximum function, you can request an expedited decision.

**For state exchanges only:** The above disclaimer applies for exigent circumstances. Expedited review may also be requested when you are undergoing a current course of treatment using a non-formulary drug.

### Demographics

Patient Information		Prescriber Information	
Patient Name:		Prescriber Name:	
DOB:	Age:	NPI#:	Specialty:
Health Plan ID#:		Phone:	Fax:
Pharmacy Name:	Pharmacy Phone:	Office Contact:	Direct Phone # or Ext:

### Medication Information

Medication: <b>Aubagio</b>	Strength: <input type="checkbox"/> 7 mg Tablet <input type="checkbox"/> 14 mg Tablet	Directions:	Quantity Dispensed:	Day Supply:
<input type="checkbox"/> New medication <input type="checkbox"/> Continuation of therapy	Start Date:	If this is continuation of therapy, please provide CHART DOCUMENTATION indicating the member showed improvement while on therapy.		

### Clinical Information

Diagnosis:	Date of Diagnosis:	
Does the member have relapsing form of Multiple Sclerosis? <span style="float: right;"><input type="checkbox"/> Yes <input type="checkbox"/> No</span> Does the member have severe hepatic impairment? <span style="float: right;"><input type="checkbox"/> Yes <input type="checkbox"/> No</span> Does the member have evidence of active infection? <span style="float: right;"><input type="checkbox"/> Yes <input type="checkbox"/> No</span> Has the member previously tried an interferon product (Avonex, Betaseron, Extavia, Rebif, Copaxone or Tecfidera)? <span style="float: right;"><input type="checkbox"/> Yes <input type="checkbox"/> No</span>		
<u>Product</u>	<u>Trial Dates</u>	<u>Reason for Failure</u>
Is the member on concomitant therapy with antineoplastic, immunosuppressive therapy, or immune modulating therapies?		<input type="checkbox"/> Yes <input type="checkbox"/> No
<u>Medication</u>	<u>Dose/Strength</u>	<u>Frequency</u>
Has the member had the following labs within the <b>past 6 months</b> ?		
Complete Blood Count (CBC)?	<input type="checkbox"/> Yes <input type="checkbox"/> No	Date: _____
Transaminase and Bilirubin level?	<input type="checkbox"/> Yes <input type="checkbox"/> No	Date: _____
PPD (tuberculin) test	<input type="checkbox"/> Positive <input type="checkbox"/> Negative	Date: _____

Member Name:	DOB:	Health Plan ID:
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Please be sure to complete and include this page with the 1<sup>st</sup> page of this form.

<b>Female Members</b>	If the member is of childbearing potential, has she had a baseline (within 1 month) negative pregnancy test prior to initiation of therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No Date: _____
	If the member is of childbearing potential, is she currently using reliable contraception during treatment? <input type="checkbox"/> Yes <input type="checkbox"/> No

**If this request is for reauthorization, please include the following documentation:**

<input type="checkbox"/> Documentation showing members disease has stabilized	<input type="checkbox"/> Documentation of no active infection
<input type="checkbox"/> Documentation that the member is <b>NOT</b> on concomitant therapy with antineoplastic, immunosuppressive, or immune modulating therapies.	<input type="checkbox"/> Documentation that the member's transaminase/bilirubin levels are being monitored consistently.

**Please provide any additional information which should be considered in the space below:**
