



Commonwealth of Massachusetts
MassHealth Drug Utilization Review Program
 P.O. Box 2586
 Worcester, MA 01613-2586

Fax: 1-877-208-7428 **Phone:** 1-800-745-7318

Strattera and Cerebral Stimulant Prior Authorization Request

MassHealth reviews requests for prior authorization (PA) on the basis of medical necessity only. If MassHealth approves the request, payment is still subject to all general conditions of MassHealth, including current member eligibility, other insurance, and program restrictions. MassHealth will notify the provider and member of its decision. Keep a copy of this form for your records. If faxing this form, please use black ink.

PA is required for Strattera (atomoxetine). In addition, PA is required for greater than 60 units per month for long-acting cerebral stimulants, greater than 90 units per month for short-/intermediate-acting cerebral stimulants, and concurrent therapy for short-/intermediate- and long acting cerebral stimulants greater than 90 units per month (all agents combined). PA is also required for any brand-name multiple-source product that has an FDA "A"-rated generic equivalent as identified by the Approved Drug Products with Therapeutic Equivalence Evaluations (also called the "Orange Book"). Additional information about Strattera and Cerebral Stimulant use can be found within the MassHealth Drug List at www.mass.gov/druglist.

Member information

Last name	First name	MI	MassHealth member ID no.	Date of birth	Sex (Circle one.) f m
Member's place of residence <input type="checkbox"/> home <input type="checkbox"/> nursing facility					

Medication information

Strattera Request <input type="checkbox"/> Strattera (atomoxetine)	Dose, frequency, and duration	Drug NDC (if known)
Cerebral Stimulant Request (Check one or all that apply.)		Dose, frequency, and duration
Long Acting <input type="checkbox"/> Adderall XR (amphetamine salts) <input type="checkbox"/> Concerta (methylphenidate) <input type="checkbox"/> Focalin XR (dexamethylphenidate) <input type="checkbox"/> Metadate CD (methylphenidate) <input type="checkbox"/> Ritalin LA (methylphenidate)	Intermediate Acting/Short Acting <input type="checkbox"/> Adderall# (amphetamine salts) <input type="checkbox"/> Focalin (dexamethylphenidate) <input type="checkbox"/> Dexedrine# (dextroamphetamine) <input type="checkbox"/> Dextrostat# (dextroamphetamine) <input type="checkbox"/> Liquadd (dextroamphetamine) <input type="checkbox"/> Metadate ER# (methylphenidate) <input type="checkbox"/> Methylin (methylphenidate) <input type="checkbox"/> Ritalin# (methylphenidate) <input type="checkbox"/> Ritalin SR# (methylphenidate)	Drug NDC (if known) Quantity requested per month Has dose consolidation been attempted? <input type="checkbox"/> Yes <input type="checkbox"/> No Please explain why not.
Indication (Check one or all that apply.) <input type="checkbox"/> Attention Deficit Hyperactivity Disorder (ADHD) <input type="checkbox"/> Narcolepsy <input type="checkbox"/> Other (Explain)		
Is member under the care of a psychiatrist or behavioral specialist? <input type="checkbox"/> Yes <input type="checkbox"/> No Name of psychiatrist or behavioral specialist _____ Telephone no.: _____ Date of last visit: _____		
Please list all medications currently prescribed for this member for this condition. _____ _____		
Please describe your new treatment plan for managing this member's condition, including discontinuation of any medications as a result of the addition of medication requested. _____ _____		

Medication information (cont.)

Please complete the following sections for Strattera requests.

Has member tried other medications in the methylphenidate class (i.e., Concerta, Focalin, Metadate, Methylin, or Ritalin) to treat this condition? Yes. Complete box A. No. Explain why not. _____

A. Drug name	Dates of use	Dose and frequency
Did member experience any of the following? <input type="checkbox"/> Adverse reaction <input type="checkbox"/> Inadequate response <input type="checkbox"/> Intolerance <input type="checkbox"/> Other		
Briefly describe details of adverse reaction, inadequate response, intolerance, or other. _____		

Has member tried other medications in the amphetamine/dextroamphetamine class (i.e., Adderall or Dexedrine) to treat this condition?

Yes. Complete box B. No. Explain why not. _____

B. Drug name	Dates of use	Dose and frequency
Did member experience any of the following? <input type="checkbox"/> Adverse reaction <input type="checkbox"/> Inadequate response <input type="checkbox"/> Intolerance <input type="checkbox"/> Other		
Briefly describe details of adverse reaction, inadequate response, intolerance, or other. _____		

Has member tried other non-stimulant medications to treat this condition?

Yes. Complete box C. No. Explain why not. _____

C. Drug name	Dates of use	Dose and frequency
Did member experience any of the following? <input type="checkbox"/> Adverse reaction <input type="checkbox"/> Inadequate response <input type="checkbox"/> Intolerance <input type="checkbox"/> Other		
Briefly describe details of adverse reaction, inadequate response, intolerance, or other. _____		

Note: You may be asked to provide supporting documentation (e.g., copies of medical records, office notes, and/or completed FDA MedWatch form).

Pharmacy information

Name	Pharmacy provider no. <i>Optional</i>	Telephone no. ()	Fax no. () <i>Optional</i>
Address		City	State Zip <i>Optional</i>

Prescriber information

Last name	First name	MI	MassHealth provider no.	DEA no.
Address			City	State Zip
E-mail address <i>Optional</i>			Telephone no. ()	Fax no. ()

Signature

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.

Prescriber's signature (Stamp not accepted.)

Date