**EMSAM (selegiline) transdermal system**
**Effective 08/01/2020**

<table>
<thead>
<tr>
<th>Plan</th>
<th>Program Type</th>
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</thead>
<tbody>
<tr>
<td>☑ MassHealth</td>
<td>☑ Prior Authorization</td>
</tr>
<tr>
<td>☑ Commercial/Exchange</td>
<td>☐ Quantity Limit</td>
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</tbody>
</table>

| Benefit | ☑ Pharmacy Benefit | ☐ Medical Benefit (NLX) |

<table>
<thead>
<tr>
<th>Specialty Limitations</th>
<th>Specialty Medications</th>
<th>Non-Specialty Medications</th>
<th>Medical Specialty Medications (NLX)</th>
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<tbody>
<tr>
<td>N/A</td>
<td>All Plans Phone: 866-814-5506 Fax: 866-249-6155</td>
<td>MassHealth Phone: 877-433-7643 Fax: 866-255-7569</td>
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<td>Commercial Phone: 800-294-5979 Fax: 888-836-0730</td>
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<td>Exchange Phone: 855-582-2022 Fax: 855-245-2134</td>
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<td>All Plans Phone: 844-345-2803 Fax: 844-851-0882</td>
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| Exceptions | N/A |

**Overview**
Emsam (selegiline transdermal system) is a monoamine oxidase inhibitor (MAOI) indicated for the treatment of adults with major depressive disorder (MDD)

**Coverage Guidelines**
Authorization may be granted for members who are currently receiving treatment with Emsam, excluding when the product is obtained as samples or via manufacturer’s patient assistance programs.

**OR**
Approval of EMSAM will be granted if the member meets the ALL the following criteria and documentation has been provided:
1. Member is at least 18 years of age **AND**
2. Member is diagnosed with major depressive disorder (MDD) **AND**
3. Member is unable to tolerate oral medications, including liquid or crushed formulations **OR**
4. Member has had a documented side effect, allergy or treatment failure with at least one SSRI and one other non-SSRI antidepressant **OR**
5. Member has a contradiction to all SSRI and non-SSRI antidepressants

**Limitations**
1. Approvals will be granted for 36 months
2. The following quantity limits apply:

<table>
<thead>
<tr>
<th>Specialty Medications</th>
<th>Quantity Limit</th>
</tr>
</thead>
<tbody>
<tr>
<td>Emsam</td>
<td>30 patches per month</td>
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</table>
Pharmacist’s Note: EMSAM®, like other antidepressants, contains a Black Box Warning:

### Suicidality and Antidepressant Drugs

Antidepressants increased the risk compared to placebo of suicidal thinking and behavior (suicidality) in children, adolescents, and young adults in short-term studies of major depressive disorder (MDD) and other psychiatric disorders. Anyone considering the use of EMSAM or any other antidepressant in a child, adolescent, or young adult must balance this risk with the clinical need. Short term studies did not show an increase in the risk of suicidality with antidepressants compared to placebo in adults beyond age 24; there was a reduction in risk with antidepressants compared to placebo in adults aged 65 and older. Depression and certain other psychiatric disorders are themselves associated with increases in the risk of suicide. Patients of all ages who are started on antidepressant therapy should be monitored appropriately and observed closely for clinical worsening, suicidality, or unusual changes in behavior. Families and caregivers should be advised for the need for close observation and communication with the prescriber. EMSAM is not approved for use in pediatric patients. Furthermore, EMSAM at any dose should not be used in children under the age of 12, even when administered with dietary modifications.

### References

1. Emsam (selegiline transdermal system) [prescribing information]. Morgantown, WV: Mylan Specialty LP; May 2020

### Review History

12/04/2006 - Implementation
11/27/2006 – Reviewed
11/26/2007 – Reviewed
01/02/2008 - Implementation
11/24/2008 - Reviewed
11/22/2010 – Reviewed
11/28/2011 – Reviewed
11/26/2012 – Reviewed
11/25/2013 – Reviewed
11/27/2017 – Reviewed
04/17/2019 – Reviewed
05/20/2020 – Reviewed and Updated May P&T Mtg; added started and stabilized statement; added QL to ‘Limitations’. Effective 8/1/20.
07/21/2021-Reviewed July P&T; no clinical changes

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