



**doxepin tablet (Silenor)**  
Effective 11/01/2022

<b>Plan</b>	<input checked="" type="checkbox"/> MassHealth <input type="checkbox"/> Commercial/Exchange	<b>Program Type</b>	<input checked="" type="checkbox"/> Prior Authorization
<b>Benefit</b>	<input checked="" type="checkbox"/> Pharmacy Benefit <input type="checkbox"/> Medical Benefit (NLX)		<input checked="" type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
<b>Specialty Limitations</b>	N/A		
<b>Contact Information</b>	<b>Specialty Medications</b>		
	All Plans	Phone: 866-814-5506	Fax: 866-249-6155
	<b>Non-Specialty Medications</b>		
	MassHealth	Phone: 877-433-7643	Fax: 866-255-7569
	Commercial	Phone: 800-294-5979	Fax: 888-836-0730
	Exchange	Phone: 855-582-2022	Fax: 855-245-2134
	<b>Medical Specialty Medications (NLX)</b>		
	All Plans	Phone: 844-345-2803	Fax: 844-851-0882
<b>Exceptions</b>	N/A		

**Overview**

Doxepin tablet is indicated for the treatment of insomnia characterized by difficulty with sleep maintenance.

**Coverage Guidelines**

Authorization may be reviewed on a case by case basis for members who are currently receiving treatment with doxepin tablet, excluding when the product is obtained as samples or via manufacturer’s patient assistance programs

**OR**

Authorization may be granted for members when all the following criteria are met, and documentation is provided:

1. Appropriate diagnosis
2. Physician documented of an inadequate response or adverse reaction to **TWO** of the following, or a contraindication to **ALL** of the following:
  - a. Belsomra or Dayvigo
  - b. doxepin capsule or liquid
  - c. eszopiclone
  - d. ramelteon
  - e. zaleplon
  - f. zolpidem immediate-release or extended release
3. **ONE** of the following:
  - a. Requested quantity is  $\leq$  1 unit/day
  - b. Medical necessity for  $>$  1 unit/day



*Additional criteria may apply for members under the age of 18. Please refer to the Pediatric Behavioral Health Medication Initiative guideline for criteria.*

### **Continuation of Therapy**

Reauthorizations requires physician documentation of continuation of therapy and positive response to therapy.

### **Limitations**

1. Initial approvals and reauthorizations will be granted for 12 months.
2. The following quantity limits apply:

doxepin tablet	30 tablets per 30 days
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### **References**

1. Silenor (doxepin) [prescribing information]. Morristown, NJ: Currax Pharmaceuticals, LLC; October 2020. Riemann D, Baglioni C, Bassetti C, et al. European guideline for the diagnosis and treatment of insomnia. *J Sleep Res.* 2017;26(6):675-700. doi:10.1111/jsr.12594
2. Sateia MJ, Buysse DJ, Krystal AD, Neubauer DN, Heald JL. Clinical practice guideline for the pharmacologic treatment of chronic insomnia in adults: an American Academy of Sleep Medicine clinical practice guideline. *J Clin Sleep Med.* 2017;13(2):307-349. doi:10.5664/jcsm.6470

### **Review History**

11/16/2022 – Created for Nov P&T; matched MH UPPL. Doxepin tablet will require a PA and QL. Criteria was matched to MH UPPL. Effective 11/1/22.

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