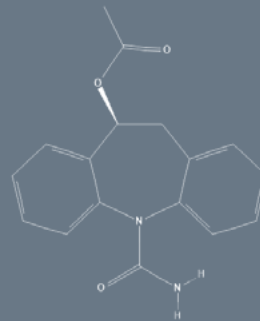


# Eslicarbazepine Acetate

## Quality Solutions

Category: Psychiatric



Authorized Distributor

USP can support your development and manufacturing activities on Eslicarbazepine Acetate-based medicines with these existing and upcoming standards.

## Official and Proposed Documentary Standards and Associated Physical Materials

### MONOGRAPH 1

<b>ESLICARBAZEPINE ACETATE</b> Published in PF 49(6) 01-Nov-2023	
USP Eslicarbazepine Acetate RS	<a href="#">USP-1249508</a>
USP Eslicarbazepine Acetate Chiral Peak ID Mixture RS	<a href="#">USP-1249519</a>
USP Eslicarbazepine Acetate Peak ID Mixture RS   <b>NEW</b>	<a href="#">USP-1249520</a>

### INCLUDED GENERAL CHAPTERS

<b>&lt;11&gt; USP REFERENCE STANDARDS</b> Official as of 01-Nov-2020
<b>&lt;197&gt; SPECTROSCOPIC IDENTIFICATION TESTS</b> Official as of 01-Sep-2021
<b>&lt;281&gt; RESIDUE ON IGNITION</b> Official as of 31-Dec-2012
<b>&lt;731&gt; LOSS ON DRYING</b> Official as of 01-Nov-2020

### INCLUDED EXCIPIENTS

<b>CROSCARMELOSE SODIUM</b> Official as of 01-May-2022	
USP Croscarmellose Sodium RS	<a href="#">USP-1150659</a>
<b>POVIDONE</b> Official as of 01-Jun-2023	
USP Povidone RS	<a href="#">USP-1551503</a>
<b>MAGNESIUM STEARATE</b> Official as of 01-Aug-2016	
USP Palmitic Acid RS	<a href="#">USP-1492007</a>
USP Stearic Acid RS	<a href="#">USP-1621008</a>
<b>SODIUM STEARYL FUMARATE</b> Official as of 01-Jun-2023	
USP Sodium Stearyl Fumarate RS	<a href="#">USP-1614705</a>
USP Stearyl Alcohol RS	<a href="#">USP-1622000</a>
USP Monostearyl Maleate RS	<a href="#">USP-1446804</a>

\* Documents in PF Online are not official and not suitable to demonstrate compliance. They may never become official.

01-May-2024. For publications related to **Eslicarbazepine Acetate** in PF online after this date, please visit the integrated [USP-NF/PF online](#).

**Disclaimer:** USP Quality Solution Sheets are provided as a convenience and for informational purposes only. Labmix24 makes reasonable efforts to provide correct information but assumes no liability for the completeness, timeliness or accuracy of the information contained in each QSS. It is the responsibility of the user to verify the information. The US Pharmacopeia remains the official source of this information.