

STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

FINGOLIMOD LAURYL SULFATE

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
FINGOLIMOD	TASCENSO ODT	48165		GPI-10	
LAURYL SULFATE				(6240702520)	

GUIDELINES FOR USE

- 1. Does the patient have a diagnosis of a relapsing form of multiple sclerosis (MS), to include clinically isolated syndrome, relapsing-remitting disease and active secondary progressive disease, and meet **ALL** of the following criteria?
 - The patient is 10 years of age or older
 - The patient had a trial of fingolimod capsules
 - The patient is unable to swallow fingolimod capsules
 - The patient had a trial of or contraindication to ONE agent indicated for the treatment of MS

If yes, continue to #2.

If no, do not approve.

DENIAL TEXT: See the denial text at the end of the guideline.

- 2. Does the patient have ANY of the following contraindications to Tascenso ODT?
 - A recent (within past 6 months) occurrence of myocardial infarction, unstable angina, stroke, transient ischemic attack, decompensated heart failure requiring hospitalization, or Class III/IV heart failure
 - A history or presence of Mobitz Type II 2nd degree or 3rd degree AV block or sick sinus syndrome, unless patient has a functioning pacemaker
 - A baseline QTc interval of 500 msec or greater
 - Current treatment with Class Ia (quinidine, procainamide, or disopyramide) or Class III antiarrhythmic drugs (amiodarone, dofetilide, dronedarone, ibutilide, or sotalol)

If yes, do not approve.

DENIAL TEXT: See the denial text at the end of the guideline.

If no, approve for 12 months by HICL or GPI-10 with a quantity limit of #1 per day.

CONTINUED ON NEXT PAGE

Copyright © 2023 MedImpact Healthcare Systems, Inc. All rights reserved. This document is proprietary to MedImpact. MedImpact maintains the sole and exclusive ownership, right, title, and interest in and to this document.

Revised: 12/21/2022 Page 1 of 3



STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

FINGOLIMOD LAURYL SULFATE

GUIDELINES FOR USE (CONTINUED)

DENIAL TEXT: *Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

Our guideline named **FINGOLIMOD LAURYL SULFATE** (**Tascenso ODT**) requires the following rule(s) be met for approval:

- A. You have a relapsing form of multiple sclerosis (a type of nerve disorder), to include clinically isolated syndrome (a type of nerve disorder that occurs once), relapsing-remitting disease (symptoms or disease returns and goes away) and active secondary progressive disease (advanced disease)
- B. You are 10 years of age or older
- C. You had a trial of fingolimod capsules
- D. You are unable to swallow fingolimod capsules
- E. You had a trial of or contraindication (harmful for) to one other agent indicated for the treatment of multiple sclerosis
- F. You do not have any of the following contraindications (harmful for) to Tascenso ODT:
 - 1. A recent (within past 6 months) occurrence of myocardial infarction (heart attack), unstable angina (chest pain), stroke, transient ischemic attack (short stroke-like attack), decompensated heart failure requiring hospitalization, or Class III/IV heart failure
 - 2. A history or presence of Mobitz Type II 2nd degree or 3rd degree AV block or sick sinus syndrome (types of irregular heartbeats), unless you have a functioning pacemaker
 - 3. A baseline QTc interval of 500 msec or greater (a measure of the speed of electrical conduction in the heart)
 - 4. Current treatment with Class Ia (quinidine, procainamide, or disopyramide) or Class III anti-arrhythmic drugs (amiodarone, dofetilide, dronedarone, ibutilide, or sotalol)

Your doctor told us [INSERT PT SPECIFIC INFO PROVIDED]. We do not have information showing you [INSERT UNMET CRITERIA]. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

CONTINUED ON NEXT PAGE

Copyright © 2023 MedImpact Healthcare Systems, Inc. All rights reserved. This document is proprietary to MedImpact. MedImpact maintains the sole and exclusive ownership, right, title, and interest in and to this document.

Revised: 12/21/2022 Page 2 of 3



STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

FINGOLIMOD LAURYL SULFATE

RATIONALE

For further information, please refer to the Prescribing Information and/or Drug Monograph for Tascenso ODT.

REFERENCES

 Tascenso ODT [Prescribing Information]. San Jose, CA: Handa Neuroscience, LLC; December 2022.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A Created: 11/22

Commercial Effective: 01/16/23 Client Approval: 12/22 P&T Approval: 10/22

Copyright © 2023 MedImpact Healthcare Systems, Inc. All rights reserved. This document is proprietary to MedImpact. MedImpact maintains the sole and exclusive ownership, right, title, and interest in and to this document.

Revised: 12/21/2022 Page 3 of 3