



New Hampshire Medicaid Fee-for-Service (FFS) Program

Prior Authorization Drug Approval Form

Encelto™ (revakinagene taroretcel-lwey)

DATE OF MEDICATION REQUEST: / /

SECTION I: PATIENT INFORMATION AND MEDICATION REQUESTED

LAST NAME:

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FIRST NAME:

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MEDICAID ID NUMBER:

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DATE OF BIRTH:

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GENDER: ☐ Male ☐ Female

Drug Name

Strength

Dosing Directions

Length of Therapy

SECTION II: PRESCRIBER INFORMATION

LAST NAME:

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FIRST NAME:

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SPECIALTY:

NPI NUMBER:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

PHONE NUMBER:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

FAX NUMBER:

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SECTION III: CLINICAL HISTORY

1. Is the patient 18 years of age and older? ☐ Yes ☐ No

2. Does the patient have a diagnosis of macular telangiectasia type 2 (MacTel) in at least one eye as evidenced by typical fluorescein leakage and at least 1 of the following? ☐ Yes ☐ No

additional features. (Select all that apply.)

- ☐ Hyperpigmentation outside a 500 micron radius from the center of the fovea
- ☐ Retinal opacification
- ☐ Crystalline deposits
- ☐ Right-angle vessels
- ☐ Inner/outer lamellar cavities

Fax to Prime Therapeutics Management LLC if medications will be dispensed by a pharmacy and will be administered by the patient or caregiver at home.

Phone: 1-866-675-7755

Fax: 1-888-603-7696

Fax to DHHS if medication is dispensed/administered by the office or outpatient setting:

Phone: 1-603-271-9384

Fax: 1-603-314-8101



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PATIENT LAST NAME:

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PATIENT FIRST NAME:

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3. Does the patient have neovascular MacTel? ☐ Yes ☐ No
4. Does the patient have evidence of advanced disease, such as significant retinal scarring or atrophy with retinal tissue that cannot be preserved? ☐ Yes ☐ No
5. Does the patient have inner segment-outer segment junction line (IS/OS) photoreceptor break and area of ellipsoid zone (EZ) loss, as measured by spectral-domain optical coherence tomography (SD-OCT), between 0.16 mm² and 2mm²? ☐ Yes ☐ No
6. Does the patient have evidence of any of the following: (select all that apply) ☐ Yes ☐ No
- ☐ Intraretinal neovascularization or subretinal revascularization (SRNV) as evidenced by hemorrhage, hard exudate, subretinal fluid, or intraretinal fluid in either eye
- ☐ Central serous chorioretinopathy in either eye
- ☐ Pathologic myopia in either eye
- ☐ Significant media or corneal opacities in either eye
- ☐ History of vitrectomy, penetrating keratoplasty, trabeculectomy, or trabeculoplasty
- ☐ Lens opacities: cortical opacity > 3 standard, posterior subcapsular opacity > standard 2, or nuclear opacity > standard 3
- ☐ Lens removal in previous 3 months or yttrium-aluminum-garnet laser treatment within 4 weeks
- ☐ History of ocular herpes virus in either eye
- ☐ Evidence of intraretinal hyperreflectivity by optical coherence tomography (OCT)
7. Does the patient have ocular or periocular infections? ☐ Yes ☐ No
8. Does the patient have a known hypersensitivity to Endothelial Serum Free Media (Endo-SFM)? ☐ Yes ☐ No
9. Prescriber attests that the patient will be monitored for signs and symptoms of vision loss and infectious endophthalmitis at baseline and periodically during treatment. ☐ Yes ☐ No
10. Prescriber attests that the patient will be monitored for signs and symptoms of retinal tears and retinal detachment. ☐ Yes ☐ No
11. Does the patient have evidence of other ocular disease that would preclude treatment of MacTel? ☐ Yes ☐ No
12. Will the patient temporarily discontinue antithrombotic medications (if applicable) prior to the insertion surgery? Leave blank if not applicable. ☐ Yes ☐ No

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13. Has the patient received intravitreal steroid therapy or intravitreal anti-vascular endothelial growth factor (VEGF) therapy for non-neovascular MacTel within the last 3 months? ☐ Yes ☐ No
14. Is there any additional information that would help in the decision-making process?
If additional space is needed, please use a separate sheet.

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: _____ **DATE:** _____

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