

Dosing frequency

Luxturna Prior Authorization Request

Your patient's benefit plan requires prior authorization for certain medications. In order to make appropriate medical necessity determinations, your patient's diagnosis and other clinical information is required. Please complete the information requested on the form below and fax this form to Priority Partners, toll-free at 1-866-212-4756 to initiate the review process. If you have questions regarding the prior authorization please contact Priority Partners at 888-819-1043 Option 4.

Patient's Name:	Date:
Patient's ID:	Patient's Date of Birth:
Physician's Name:	
Specialty:	NPI#:
Physician Office Telephone:	Physician Office Fax:
<u>Referring</u> Provider Info:	equesting Provider
Name:	NPI#:
Fax:	Phone:
<u>Rendering</u> Provider Info: 🗖 Same as R	eferring Provider 🖵 Same as Requesting Provider
Name:	NPI#:
Fax:	Phone:
	t to dosing limits in accordance with FDA-approved labeling, pendia, and/or evidence-based practice guidelines.
Required Demographic Information:	
Patient Weight:	ka

Patient Weight:	kg	
Patient Height:	<i>cm</i>	
Please indicate the place of service for th Ambulatory Surgical On Campus Outpatient Hospital	te requested drug: ☐ Home ☐ Office	□ Off Campus Outpatient Hospital
Drug Information:		
Strength/Measure		$_Units \square ml \square Gm \square mg \square ea \square Un$
Directions(sig)		_Route of administration

Send completed form to: Priority Partners Fax: 1-866-212-4756

Note: This fax may contain medical information that is privileged and confidential and is solely for the use of individuals named above. If you are not the intended recipient you hereby are advised that any dissemination, distribution, or copying of this communication is prohibited. If you have received the fax in error, please immediately notify the sender by telephone and destroy the original fax message. JHHC Luxturna SGM 2458-A – 07/2023.

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Criteria Questions:

What is the ICD-10 code?

1. What is the diagnosis?

□ Biallelic RPE65 mutation-associated retinal dystrophy (If checked, go to 2)

□ Other, please specify. _____ (*If checked, go to 2*)

2. Is there confirmation of bi-allelic pathogenic and/or likely pathogenic RPE65 gene mutations?

 \square Yes, Continue to 3

□ No, Continue to 3

3. Please indicate which of the following genetic tests was performed to confirm bi-allelic pathogenic and/or likely pathogenic RPE65 gene mutations. *ACTION REQUIRED*: Attach genetic test results (single gene test or multi gene panel test) confirming a genetic diagnosis of pathogenic/likely pathogenic biallelic RPE65 gene mutations.

□ Single gene panel test (*If checked, go to 4*)

□ Multi gene panel test (If checked, go to 4)

□ None of the above (*If checked, go to 4*)

4. Are the RPE65 gene mutations classifications based on the current American College of Medical Genetics and Genomics (ACMG) standards and guidelines for the interpretation of sequence variants?

□ Yes, *Continue to 5* □ No, *Continue to 5*

5. Please provide the date of the genetic test.

Date: ______MM/DD/YY (If checked, go to 6)

 $\Box \text{ Unknown} (If checked, go to 6)$

6. Has pathogenicity of the RPE65 mutations been affirmed within the last 12 months?

□ Yes, Continue to 7

□ No, Continue to 7

7. What is the patient's age?

Less than 12 months of age (*If checked, go to 8*)

 \Box 12 months to 64 years of age (*If checked*, go to 8)

 \square 65 years of age or older (*If checked, go to 8*)

8. Which of the following test(s) was performed to confirm that the patient has viable retinal cells in each eye to be treated?

□ Optical coherence tomography (OCT) (*If checked, go to 9*)

□ Ophthalmoscopy (*If checked, go to 9*)

□ Optical coherence tomography (OCT) and ophthalmoscopy (*If checked, go to 9*)

□ None of the above (If checked, go to 9

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9. Does the patient have an area of the retina within the posterior pole of greater than 100 micrometer thickness shown on optical coherence tomography (OCT)?

 \square Yes (If checked, go to 12)

 \square No (If checked, go to 10)

 \Box Unknown (*If checked, go to 10*)

10. Within the posterior pole, how many disc areas of the retina are without atrophy or pigmentary degeneration?

 \square 3 or more (*If checked, go to 12*)

 \Box Less than 3 (*If checked, go to 11*)

Unknown (If checked, go to 11)

11. Is the patient's remaining visual field within 30 degrees of fixation as measured by a III4e isopter or equivalent?

 \square Yes (If checked, go to 12)

 \square No (If checked, go to 12)

 $\Box \text{ Unknown } (If checked, go to 12)$

12. Has the patient had the requested drug in the past?

□ Yes, Continue to 13

□ No, *No Further Questions*

13. Please select the eye which was treated in the past.

□ Right eye (*If checked, go to 14*)

Left eye (If checked, go to 15)
Both eyes (If checked, no further questions)

14. Is this request for a right eye treatment?

□ Yes, right eye (If checked, no further questions)

□ No, left eye (*If checked, no further questions*)

15. Is this request for a left eye treatment?

□ Yes, left eye (*If checked, no further questions*)

□ No, right eye (*If checked, no further questions*)

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by Priority Partners.

Х

Prescriber or Authorized Signature

Date (mm/dd/yy)

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