

## **Oxlumo**

## **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 along with supporting clinical documentation 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:	
Referring Provider Info: ☐ Same as Re	equesting Provide	er	
Name:		NPI#:	
Fax:		Phone:	
Rendering Provider Info: ☐ Same as Re	eferring Provide		
Name:		· •	
Fax:		Phone:	
Required Demographic Information:  Patient Weight:	kg		
Patient Height:	cm		
Please indicate the place of service for the	requested drug:		
	☐Home	☐ Off Campus Outpatient Hospital	
On Campus Outpatient Hospital	<b>□</b> Office		
Drug Information:			
Strength/Measure		Units □ ml □ Gm □ mg □ ea □ Un	
		_	
Directions(sig)		Route of administration	
Dosing frequency		- · · · · · · · · · · · · · · · · · · ·	

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XPrescriber or Authorized Signature	Date (mm/dd/yy)
I attest that this information is accurate and true, and that doci information is available for review if requested by Priority Pari	
☐ Yes, No Further Questions ☐ No, No Further Questions ☐ Unknown, No Further Questions	
☐ Yes, Continue to 4 ☐ No, No Further Questions  4. Has the patient's urinary and/or plasma oxalate decreased or	
2. Does the patient have a documented diagnosis of primary by the following? <i>ACTION REQUIRED</i> : If Yes, attach supporting a mutation in the alanine: glyoxylate aminotransferase (AGXT) absent or significantly reduced alanine: glyoxylate aminotransferase demonstrating absent or significantly reduced alanine: glyoxylate aminotransferase demonstrating absent or significantly reduced alanine: glyoxylate   ☐ Yes, <i>Continue to 3</i> ☐ No, <i>Continue to 3</i> 3. Is the patient currently receiving treatment with the requested	g chart note(s) for molecular genetic tests showing gene or a liver enzyme analysis demonstrating erase (AGT) activity. a) Molecular genetic test (AGXT) gene, b) Liver enzyme analysis te aminotransferase (AGT) activity.
1. What is the diagnosis?  ☐ Primary hyperoxaluria type (PH1), Continue to 2  ☐ Other, please specify, Conti	nue to 2
What is the ICD-10 code?	
Criteria Questions:	

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

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