

Start Form



> Before submitting the Start Form to Alnylam Assist®, patient **and** prescriber signatures are required

> Patients currently prescribed an Alnylam medicine who are enrolled in Alnylam Assist do not need to complete Sections 1 - 4

For Patients

Alnylam Assist® Enrollment

(Sections 1 – 4 to be read and completed by Patient or Patient's Authorized Representative)

The purpose of this form is to permit Alnylam Assist® participants to receive additional information and support ("Patient Support") from Alnylam Pharmaceuticals, Inc., its affiliates, representatives, agents, and contractors ("Alnylam"). Alnylam Assist® provides Patient Support to eligible patients who have been prescribed an Alnylam medicine. This includes: (1) providing reimbursement and financial support to eligible patients (such as investigating your insurance coverage, confirming out-of-pocket costs, and reviewing eligibility for financial assistance); (2) working with you and your provider to fill your prescription; (3) providing you with disease and medication-related educational resources and communications; and (4) contacting you to participate in disease and medication-related market research panels or surveys. Your authorization in this form will relate to information and support with respect to any Alnylam medicine you have been prescribed or may be prescribed in the future.

Please read this form carefully and ask any questions that you may have before signing.

1. Patient Information									
Name (First, MI, Last):									
Date of Birth: Month/Day/Year					Email:				
Street Address:									
City:		Sta	te:						
Home Phone #: Preferred Okay to leave message	Mobile Phone #: □F	obile Phone #: Preferred Okay to leave message			Alternative Phone # (if available): Preferred Okay to leave message				
Caregiver Name (optional):	Caregiver Relationship to Patient (optional):		Caregiver Phone (optional): Okay to leave message						
Caregiver Email (optional):				ge translation? Yes, translation needed No If yes, please indicate language:					
2. Insurance Information Attac	h a co	py of both sides of	your INSI	URANCE and PRES	CRIP	TION cards 🔲 Che	ck if you d	o not have insurance	
Primary Insurance Provider:	Emp	oloyer Name:		Policy Number:			Group Number:		
Policyholder Name (First, MI, Last), if other than the patient:				Policyholder Date of Birth: Month/Day/Year			Insurance Phone:		
Pharmacy Plan Provider (if applicable):	Poli	olicy Number:		Group Number:		Rx Bin Number:		Rx PCN Number:	
Policyholder Name (First, MI, Last), if other than the patient:				Policyholder Date of Birth: Month/Day/Year			Insurance Phone:		
Secondary Insurance Provider (if applicable):	Emp	bloyer Name:		Policy Number:			Group Number:		
Policyholder Name (First, MI, Last), if other than the patient:				Policyholder Date of Birth: Month/Day/Year Insurance Phone				e Phone:	



Print Patient or Authorized Patient Representative Name

Start Form



3. Authorization to Share Protected Health Information

By signing below, I authorize my healthcare providers, including my physicians and pharmacies ("My Providers") and my health insurance plan ("My Plan") to share my medical information (such as information about my diagnosis, prescriptions, and treatment) and my insurance information ("My Information") with Alnylam so that Alnylam can provide Patient Support. I authorize My Providers to use My Information to provide me with certain offerings related to my treatment and any Alnylam medicine My Providers may prescribe for me at any time. I understand that my pharmacy will receive payment from Alnylam for disclosing My Information to Alnylam. I understand that once My Information has been disclosed, federal privacy laws may no longer protect the information. However, I understand that Alnylam agrees to protect My Information by using and disclosing it only for purposes described in this Authorization or as required by law. I understand that I may refuse to sign this Authorization, and that my treatment, insurance enrollment, and eligibility for insurance benefits are not conditioned upon signing this Authorization. I also understand, however, that refusing to sign this Authorization means that I may not participate in Alnylam Assist® and may not be able to take advantage of other offerings by Alnylam. I may cancel or revoke this Authorization at any time by mailing a letter to Privacy Officer at Alnylam, Attn: Legal Department, 675 West Kendall Street, Cambridge, MA 02142 or by sending an email to privacy@alnylam.com. I understand that if I revoke this Authorization, My Providers and Alnylam will stop using and sharing My Information under this Authorization, but my revocation will not affect uses and disclosures of My Information prior to my revocation in reliance upon this Authorization.

This Authorization expires ten (10) years from the date signed below, or earlier if required by state or local law, unless I revoke it before then. I understand that I may receive a copy of this Authorization.

For information about how your personal data are processed as a part of our program, please visit www.alnylampolicies.com/privacy.

Signature of Patient or Authorized Patient Representative

Relationship to Patient	Date					
4. Authorization for Alnylam Assist® and Commun By signing below, I confirm I would like to enroll in the Aln Patient Support. I understand that Alnylam Assist® is an o	ylam Assist® program and authorize Alnylam to provide me with					
the Patient Support, administering the Alnylam Assist® proobligations. For example, Alnylam may communicate with caregiver, use My Information to tailor the Alnylam Assist®-with My Providers about dispensing Alnylam medicine to combine it with information about other patients, and use I understand that the administration of the program migh	with My Providers or My Plan in connection with providing ogram, or as otherwise required by Alnylam to meet its legal in me (such as by mail, phone, email, and/or text message) or my related communications to my needs, and share information me. I understand that Alnylam may de-identify My Information, e the resulting information for Alnylam's business purposes. It involve the use of artificial intelligence technologies to party vendors might de-identify My Information for machine					
	X					
Print Patient or Authorized Patient Representative Name	Signature of Patient or Authorized Patient Representative					

Relationship to Patient

Date



Start Form



For Healthcare Providers

(Sections 5 – 7 to be read and completed by **Healthcare Provider**)

5. Prescribe	er Inform	ation											
						/Clinic/Institution Name: Specialty:							
Practice Street Address: City						City:			State:	ZIP:			
Phone:	F	x: Tax ID #: National P				nal Provide	rovider ID (NPI) #: Si			State License #:			
Office Contact Name: Phone:						e:	E			Email:			
Referring Physicia	n:												
☐ Specialty Dist	irmacy: PANTHERx tributor (McKe	○ No preference esson Specialty or		_			Inknow			' 	d First Treatment I	Date:	
6. OXLUMO® Full Patient Name			ption (This	is a prescription;	; a pre	scriber's si	ignatur	e and date			sirth: Month/Day/\	/ear:	
Primary Diagnosis	,												
, ,	Treatment	Patient Weight (in kg)	Date Patient Weight Taken	Body Weight	t	OXLUM	10 Prescription			otal ted Dose	Number of Vials/Treatment	Refills	
OXLUMO Starting Dose (given at 1-month intervals)		e			10 kg to less than 20 kg 6 mg/kg once monthl		nly for 3 doses	(mg)		94.5 mg/0.5 mL vial(s)	Refill x 2		
subcutaneous use, 94.5 mg/0.5 mL	Ongoing Do: (begin 1 month after the last starting dose)	se		Less than 10 kg 10 kg to less than 20 kg and above	20 kg	3 mg/kg o	kg once every 3 months		(mg) (mL)		94.5 mg/0.5 mL vial(s)	Refill x 8 Refill x 2 Other	
Any known allergie	es? 🗌 Yes 🛭	No If yes, plea	ase list:										
List or attach a list	of concomita	nt medications:											
Special Instruction	ns:												
☐ If acquiring thro	ough Orsini or	PANTHERx, please	check here to a	authorize ancillary	supp	lies, such as	s needle	es and syrin	ges, as ne	eeded to	administer treatm	nent.	
☐ I confirm that my	y patient is bein	g prescribed OXLUM	O for the treatm	ent of primary hype	roxalu	ria type 1 (PI	H1) to lo	wer urinary a	and plasm	na oxalate	levels in children a	nd adults.	
		y behalf for the lim cific prescription re									ε.		
X													
Prescriber Signatu	ıre (No Stamps	s) Dispense as Writt	en					Date					
X													
Prescriber Signatu	ıre (No Stamps	s) Substitution Perr	nitted					Date					
Desired Site of Ca ☐ Home Injection ☐ Alternate Medic Facility Name/Add	n (see patient h cal Facility (pro	nome address) ovide facility name	and address)					office addre		home)			





7. Prescriber Declaration

By signing below, I certify that:

- > The information contained in this form is complete and accurate to the best of my knowledge
- ▷ I understand that Alnylam is not responsible for filing claims or submitting other information to my patient's insurer and that the information provided by Alnylam Assist® is educational in nature
- ▷ I understand that my patient may authorize Alnylam Assist® to provide Patient Support. I understand that this program does not include individual treatment or medical advice to the patient, and it does not replace the medical treatment and care provided by me as the patient's healthcare provider
- ▷ I further certify that I understand that any support provided by Alnylam Assist® on behalf of any patient is not made in exchange for any express or implied agreement or understanding that I would recommend, prescribe, or use OXLUMO® (lumasiran) or any other Alnylam product, and any decision to prescribe OXLUMO was, and in the future will be, based solely on my determination of medical necessity
- ▷ I have obtained the required authorizations from my patient to release the referenced medical and/or other patient information relating to my patient's treatment to Alnylam Assist®

x	
Prescriber signature (stamps not acceptable)	Date

INDICATION

OXLUMO® (lumasiran) is indicated for the treatment of primary hyperoxaluria type 1 (PH1) to lower urinary and plasma oxalate levels in children and adults.

IMPORTANT SAFETY INFORMATION

Adverse Reactions

The most common (≥20%) adverse reaction reported in patients treated with OXLUMO was injection site reaction. Injection site reactions included erythema, swelling, pain, hematoma, pruritus, and discoloration.

Pregnancy and Lactation

No data are available on the use of OXLUMO in pregnant women. No data are available on the presence of OXLUMO in human milk or its effects on breastfed infants or milk production. Consider the developmental and health benefits of breastfeeding along with the mother's clinical need for OXLUMO and any potential adverse effects on the breastfed child from OXLUMO or the underlying maternal condition.

For additional information about OXLUMO, please see full Prescribing Information.



Fax the completed Start Form

Call Alnylam Assist® at 1-833-256-2748 8AM-6PM, Monday-Friday

For more information, visit www.AlnylamAssist.com