PROLASTIN-C LIQUID Prescription and Enrollment Form Fax completed form to: 1-866-588-6940







PATIENT INFORMAT	TION											
Patient first name	Middle in	nitialLast n	ame		s	SN (last 4 digits	only)	DOB_	//_	Gender _]F	
Address				City_			State			Zip		
Best contact number		Home	Mobile [Work Emai	il address_							
INSURANCE INFOR	MATION Please attach a c	opy of both	sides of all	patient's med	dical and	l prescriptio						
Primary incurance				Secondar	v incuranc	۵	_	heck if pa	tient does	not have ins	surance	
Primary insurance Insurance contact number												
Policy ID#												
Group ID#												
Policy holder full name												
Relationship to patient												
tologionomp to patient _					p to patie	<u></u>						
	Steps to e-Prescribe PROLASTIN-C LIQUID	 Prescribe P Choose EV 	ROLASTIN-C ERSANA and	C LIQUID prescri LIQUID or EVERSANA I	LIFE SCIE	NCE SERVICES	S as the dispe	nsing phari	пасу			
MEDICAL INFORMA	TION Please include a	copy of patie	ent's clinica	al notes								
	Alpha₁-Antitrypsin Deficiency E8				3.1	Other						
										% pro	edicted	
Serum AAT Level ma/dL or u.M. Allergies \(\sumbox{None or } \sumbox{Specify} \)												
Smoking History												
Medical History COPD Asthma Emphysema Other												
_					access	Peripheral	Central]Port		_/		
PROLASTIN-C LIQU	IID PRESCRIPTION INFORMA	TION										
Dose Directions					Quantity/Refills							
60 mg/kg (+/- 10%) I\	· ·	Rate: As tolerated by patient up to 0.08 mL/kg/min			_	Dispense up to 28-day supply. Refill x1 year unless otherwise noted						
Other dose/frequency	Other rate				_ Other							
	☐ lb ☐ kg											
Epinephrine injection	n into the muscle from auto-device a	s needed for se	vere allergic re	eaction or anaph	ylaxis							
EMLA or lidocaine 2.	.5%/prilocaine 2.5%: Apply 2.5 g over	er 20-25 cm ² of s	skin surface at	t least 1 hour pri	or to punct	ure						
Premedication/other	orders:											
	Peripheral IV line				Implanted port	/central line (each lumei	7)				
Intravenous access and flush orders:	☐ Normal saline: 3-5 mL before infusion and 3-5 mL after in				nfusion Normal saline: 5-10 mL before infusion and 5-10 mL after infusion						on	
and nush orders:	☐ Other				Heparin: 100 units/mL, 5 mL after infusion, and (frequency)						ency)	
First influsion leastion are	eference: Home or Medica											
		arracility (name,	priorie oi prei	еней тасшіў, іг а	iriy)							
First infusion in home nu	irsing orders: line with 250-500 mL of normal salin	🗆 0#		-+ 10.10		ta :=6i.a.						
	at KVO	Provide infusion supplies, including syringes and needles, to safely administer prescribed medication.										
Monitor patient including VS before, Q15 during, and 30 minutes post infusion							ricculco, to ot	nery admin	Stor preser	ibca medicati		
. ,	-50 mg IV as directed by nurse for a	llergic reaction				Ì						
PRESCRIBER INFO												
Prescriber first name	iber first name Prescriber last nam											
					StateZip							
Office contact name Office contact phone Office contact email address												
Office contact email add	ress											
knowledge. I also atte	uthorize this prescription and certify est that I have obtained the patient's A, and/or their agents. If the patient i	authorization to	release the a	bove information	n and such	other personal	information as	may be no	ecessary to		1	
Dragaribar Ciar	a a truma						D-4-					

Dispense as Written

Substitution Permitted

Fax completed form and documentation to 1-866-588-6940. To reach the PROLASTIN DIRECT team, call 1-800-305-7881.

PRIOR AUTHORIZATION CHECKLIST

Please note that the information listed below outlines what is typically required for insurance to review the patient's eligibility. If any of the following information is not provided, it may delay approval or be cause for a denial.

NEW DIAGNOSIS OF AATD

Required documentation for insurance review

(A) Laboratory work	(B) Most recent clinical and diagnostic test results Documenting history of Emphysema					
 □ AAT serum concentration Most major insurance policies define acceptable levels as: ≤11 μM (11 μmol/L) or 80 mg/dL by radial immunodiffusion or <50 mg/dL if measured by nephelometry 	 Patient's medical records demonstrating diagnosis of AATD and clinical evidence of emphysema/worsening of emphysema due to lung disease exacerbations 					
☐ Phenotype or Genotype	☐ Diagnostic imaging-chest X-ray, CT scan					
PiZZ PiZ (null) Pi (null, null)	 Evidence of lung function decline- forced expiratory volume (FEV) and pulmonary function test (PFT) 					
PiSZ	☐ Patients clinical notes					
Other: One-on-one discussion may be required with the insurance plan medical director	☐ Smoking history					
ay be required by insurance plan for approval						
Supplemental documentation						
☐ Letter of Medical Necessity						
☐ Peer-reviewed articles supporting diagnosis and treatment						
☐ IgA antibody results (may be required for certain insurance plan approvals)						

IMPORTANT SAFETY INFORMATION

PROLASTIN®-C LIQUID is an alpha,-proteinase inhibitor (human) (alpha,-PI) indicated for chronic augmentation and maintenance therapy in adults with clinical evidence of emphysema due to severe hereditary deficiency of alpha,-PI (alpha,-antitrypsin deficiency).

Limitations of Use

- The effect of augmentation therapy with any alpha,-PI, including PROLASTIN-C LIQUID, on pulmonary exacerbations and on the progression of emphysema in alpha,-PI deficiency has not been conclusively demonstrated in randomized, controlled clinical trials
- · Clinical data demonstrating the long-term effects of chronic augmentation or maintenance therapy with PROLASTIN-C LIQUID are not available
- PROLASTIN-C LIQUID is not indicated as therapy for lung disease in patients in whom severe alpha, -PI deficiency has not been established

PROLASTIN-C LIQUID is contraindicated in immunoglobulin A (IgA)-deficient patients with antibodies against IgA or patients with a history of anaphylaxis or other severe systemic reaction to alpha₁-PI products.

Hypersensitivity reactions, including anaphylaxis, may occur. Monitor vital signs and observe the patient carefully throughout the infusion. If hypersensitivity symptoms occur, promptly stop PROLASTIN-C LIQUID infusion and begin appropriate therapy.

Because PROLASTIN-C LIQUID is made from human plasma, it may carry a risk of transmitting infectious agents, eg, viruses, the variant Creutzfeldt-Jakob disease (vCJD) agent, and, theoretically, the Creutzfeldt-Jakob disease (CJD) agent. This also applies to unknown or emerging viruses and other pathogens.

The most common adverse reactions during PROLASTIN-C LIQUID clinical trials in >5% of subjects were diarrhea and fatigue, each of which occurred in 2 subjects (6%).

Please see accompanying full Prescribing Information for PROLASTIN-C LIQUID.