Pulmonary Arterial Hypertension (PAH) Infused/Inhaled Enrollment Form



 Fax Referral To: 1-855-297-1270
 Phone: 1-888-280-1190

 Address: 6020 Ave Roberto Sanchez Vilella Carolina, PR 00982
 NCPDP: 4026325

Six Simple Steps	s to Submitting a Referral	
PATIENT INFORMATION (Complete or include demograph	nic sheet)	
Patient Name:		
Address:		
Preferred Contact Methods: Phone (to primary # provided below)	below) Text (to cell # pro	vided below) 🗌 Email (to email provided
Note: Carrier charges may apply. By providing the phone numb	per(s) and email address abov	e, you are consenting to receive
automated calls, emails and/or text messages from CVS Speci	alty® about your prescription	(s), account, and health care. Standard data
rates apply. Message frequency varies. If unable to contact via		
Primary Phone:		
		Primary Language:
Parent/Caregiver/Legal Guardian Name (Last, First):	Relationship to p	Datient:
PRESCRIBER INFORMATION		
Prescriber's Name:		
NPI #: DEA #: Group or Hospital:		
Address: C	City, State, ZIP Code:	
Phone: Fax Conta	ct Person: C	ontact's Phone:
3 INSURANCE INFORMATION Please fax copy of prescription	n and insurance cards with this fo	orm, if available (front and back)
4 DIAGNOSIS AND CLINICAL INFORMATION		
	Office Other:	
Diagnosis (ICD-10):		
Date of Diagnosis:		
	I27.20 Pulmonary Hyperte	-
		nolic Pulmonary Hypertension
	I27.89 Other Specified Puli	nonary Disease
Other Code: Description		
Patient Clinical Information:		
New York Heart Association (NYHA) Functional Classification:		
6 Minute Walk Distance: meters		
Is patient currently on another therapy for pulmonary hyperter	nsion? 🗌 Yes 🗌 No	
If Yes, name of drug(s):		
Weight: lb/kg Height: in/cm Allergies:		
Attach copies of: History and Physical Right Heart Cathe		el Blocker Statement 🗌 Echocardiogram
Nursing: Not Needed Pre-hospital/Pre-home Teaching		
		arsing Follow-up
Start of care date: Number of visits:	—	
Prostacyclin Referral Information:		
Check the boxes below to designate which items are includ		
PAH diagnosis and ICD-10 code (designated on PAH referral fo		
Is Medicare Part B the primary insurance for this referral?	L No	
Current H&P (within 6 months); Date of H&P:		
Right Heart Catheterization (RHC); Check below if included i	•	
Mean PA Pressure (or systolic/diastolic) > 25 mmHg at re	est or > 30 mmHg with exertion	
Cardiac Output		
	llary Wedge Pressure (or LVED	P) < 15 mmHg
Echocardiogram		
Calcium Channel Blocker statement with supporting docum		
Patients with the following disease states will require docum		
heart disease, valvular heart disease, lung disease, sarcoidosis	and other co-morbidities, exce	pt for the ones listed in WHO Group I
category		

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	Tyvaso,	Tyvaso DPI, Epopros	tenol (Generic Flolan)		
	Please Con	nplete Patient and	Prescriber Information		
Patient Name:		Patient DOB:	Patient Phone:		
Patient Address:					
		Prescriber Phone:			
5 PRESCRIPTION	N INFORMATION				
INHALED PRODUC	<u>:TS:</u>				
MEDICATION	STRENGTH		DOSE & DIRECTIONS	QUANTITY/REFILLS	
Tyvaso (treprostinil) Inhalation Solution	Tyvaso Inhalation System Starter Kit Tyvaso Refill Kit	 Start with 3 breaths (18 mcg) four times daily. Increase by 3-4 breaths at 1-2 week intervals, if tolerated, until the target dose of 9 breaths (54 mcg) four times daily. Other: 		Quantity: 28-day supply Refills:	
Tyvaso DPI (Treprostinil)	Tyvaso DPI Titration Kit 16 mcg/32 mcg 16 mcg/32 mcg/48 mcg Tyvaso DPI Maintenance Kit 16 mcg 32 mcg 48 mcg 64 mcg 80 mcg: 32 mcg/48 mcg	Target dose: 48 mcg 64 mcg Other mcg per treatment session, 4 times daily Start with one 16 mcg cartridge per treatment session, 4 times daily. Increase cartridge strength by 16 mcg per treatment session every week as tolerated to selected target dose. Inhale one breath per cartridge 4 times daily Other:		 Tyvaso DPI Titration Kit Quantity: 28-day supply Refills: 0 Tyvaso DPI Maintenance Kit Quantity: 28-day supply Refills: 	
Patient is interested in pa		SIGNATURE NOT ALLOWED	, , , , , , , , , , , , , , , , , , , ,	ed as needed for administration	
DAW / May Not Substitu	Brand Medically Necessary / Do Not Sub	stitute / No Substitution /	May Substitute / Product Selection Permitted / Substitution Permissible Prescriber's Signature:	Date:	
CA. MA. NC & PR: Interd	change is mandated unless Prescriber writes	the words " No Substitution "	ATTN: New York and Iowa providers, pl	ease submit electronic prescription	

The information provided above is true and accurate to the best of my knowledge, with supporting documentation in the patient's medical record. By signing above, I hereby authorize CVS Specialty Pharmacy and/or its affiliate pharmacies to complete and submit prior authorization (PA) requests to payors for the prescribed medication for this patient and to attach this Enrollment Form to the PA request as my signature.

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	Please	Complete Patient and P	Prescriber Information	
		Patient DOB:	Patient Phone:	
			Prescriber Phone:	
INFUSED THERAF	<u>STRENGTH</u>	DO	SE & DIRECTIONS QL	JANTITY/REFILLS
MEDICATION	STRENGTH	SC continuous over 24 ho		JANTITY REFILLS
Remodulin (treprostinil) for injection	☐ 1 mg/mL, 20 mL vial ☐ 2.5 mg/mL, 20 mL vial ☐ 5 mg/mL, 20 mL vial ☐ 10 mg/mL, 20 mL vial	Initial dose: ng/kg/ days until goal of ng/kg/ Change infusion site every Palliative med PRN Pump: 2 CADD-MS3 pumps* IV infusion continuous over Initial dose: ng/kg/ days until goal of ng/kg/ days until goal of ng/kg/ Diluent: Check one (Sterile di checked) 0.9% NaCl for injection Epoprostenol Sterile dilue Pump: 2 CADD-Legacy Pumps [2 CADD-MS 3 Pumps* CVC Care:	min. Titrate byng/kg/min every g/kg/min achieved. days. *For pediatric or low weight patients ONLY er 24 hours min. Titrate byng/kg/min every g/kg/min achieved. luent for Remodulin will be used if no box is Sterile Water for injection mtSterile diluent for Remodulin	Quantity: One-month supply of drug and supplies. Dosing weight: kg/lb Refills:
☐ Treprostinil (Generic Remodulin)	☐ 1 mg/mL, 20 mL vial ☐ 2.5 mg/mL, 20 mL vial ☐ 5 mg/mL, 20 mL vial ☐ 10 mg/mL, 20 mL vial	days until goal of n <u>Diluent</u> : Check one (Sterile di checked) 0.9% NaCl for injection Epoprostenol Sterile dilue <u>Pump</u> : 2 CADD-Legacy Po <u>CVC Care:</u>	min. Titrate byng/kg/min every g/kg/min achieved. luent for Treprostinil will be used if no box is	Quantity: One-month supply of drug and supplies. Dosing weight: kg/lb Refills:
Veletri (epoprostenol) for injection	☐ 0.5 mg vial ☐ 1.5 mg vial	IV infusion continuous over 24 hours Initial dose:ng/kg/min. Titrate byng/kg/min every days until goal ofng/kg/min achieved. Discharge dose:ng/kg/min Concentration:ng/mL Diluent: Check one (0.9% Sodium Chloride will be used if no box is checked) I 0.9% NaCl for injection Sterile Water for injection Pump: I 2 CADD-Legacy Pumps I 2-CADD Solis Pumps CVC Care: I Dressing change every days. Per IV standard of care		Quantity: 30-day supply of drug and supplies. Dosing weight: kg/lb Refills:
Epoprostenol (Generic Veletri)	☐ 0.5 mg vial ☐ 1.5 mg vial	IV infusion continuous over 24 hours Initial dose: ng/kg/min. Titrate byng/kg/min every days until goal of ng/kg/min achieved. Discharge dose: ng/kg/min Concentration: ng/mL <u>Diluent:</u> Check one (0.9% Sodium Chloride will be used if no box is checked) 0.9% NaCl for injection Sterile Water for injection <u>Pump:</u> 2 CADD-Legacy Pumps 2-CADD Solis Pumps <u>CVC Care</u> : days. Per IV standard of care		Quantity: 30-day supply of drug and supplies. Dosing weight: kg/lb Refills:
Patient is interested in p		STAMP SIGNATURE NOT ALLOWED	Ancillary supplies and kits provided as neede	d for administration
			AMP SIGNATURE NOT ALLOWED)	
"Dispense As Written" DAW / May Not Substi	' / Brand Medically Necessary / Do N itute	lot Substitute / No Substitution /	May Substitute / Product Selection Permitted / Substitution Permissible	
Prescriber's Signature:Date:Date:		Prescriber's Signature:	Date:	

to complete and submit prior authorization (PA) requests to payors for the prescribed medication for this patient and to attach this Enrollment Form to the PA request as my signature. CONFIDENTIALITY NOTICE: This communication and any attachments may contain confidential and/or privileged information for the use of the designated recipients named above. If you are not the intended recipient, you are hereby notified that you have received this communication in error and that any review, disclosure, dissemination, distribution or copying of it or its contents is prohibited. If you have received this communication in error, please notify the sender immediately by telephone and destroy all copies of this

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