Pulmonary Arterial Hypertension (PAH) Infused/Inhaled/Injectable 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

PATIENT INFORMATION (Com	-	raphic sheet)	ganororrai	
			OOB:	Gender: 🗌 Male 📗 Female
	ne (to primary # provid	ded below) 🗌 Text	(to cell # provide	ed below) 🗌 Email (to email provided
below)				
Note: Carrier charges may apply. By				
				account, and health care. Standard data
rates apply. Message frequency vari		-	-	y will attempt to contact by phone.
				mary Language:
				ient:
2 PRESCRIBER INFORMATION				
Prescriber's Name:		State License #:		
NPI #: DEA #:				
Address:				
				act's Phone:
3 INSURANCE INFORMATION F				
		ption and insurance ca	arus with this form	, ii avaliable (ii oi it ai iu back)
4 DIAGNOSIS AND CLINICAL II		. 🗆		
Needs by Date:	Ship to: [_] Pati	ient 🔛 Office 🔛 Of	ther:	
Diagnosis (ICD-10):				
Date of Diagnosis:				
☐ I27.0 Primary Pulmonary Hypert	ension	🗌 I27.20 Pulmoi	nary Hypertensi	on, Unspecified
☐ I27.21 Secondary Pulmonary Arte	erial Hypertension	I27.24 Chroni	c Thromboemol	ic Pulmonary Hypertension
☐ I27.83 Eisenmenger's Syndrome)	☐ I27.89 Other \$	Specified Pulmo	nary Disease
☐ Other Code:	Description			
Patient Clinical Information:				
New York Heart Association (NYHA)) Functional Classificati	ion: 🔲 I 🗎 II 🦳	III 🗆 IV	
6 Minute Walk Distance:			_	
Is patient currently on another thera		ertension? Yes	□No	
If Yes, name of drug(s):				
Weight:lb/kg Height: _		ies.		
				locker Statement
Nursing: Not Needed Pre-hos	•	ng 🔲 In-hospital Te	aching Nursi	ng Follow-up
Start of care date: I	lumber of visits:			
Prostacyclin Referral Information	<u>1</u>			
Check the boxes below to designate	te which items are inc	cluded in this fax:		
PAH diagnosis and ICD-10 code (d	esignated on PAH referr	al form)		
Is Medicare Part B the primary insurar	nce for this referral? \Box `	Yes 🗌 No		
Clinical documentation				
Current H&P (within 6 months);	Date of H&P:			
Right Heart Catheterization (RHC		· ·		
Mean PA Pressure (or systolic			with exertion	
Cardiac Output	Cardiac Inde			
Pulmonary Vascular Resistan	ce	Capillary Wedge Pres	sure (or LVEDP)	< 15 mmHg
Echocardiogram				
Calcium Channel Blocker staten	• • • • •			
				roportion with the secondary disease: Left
heart disease, valvular heart diseas	e, lung disease, sarcoido	osis and other co-mo	rbidities, except	or the ones listed in WHO Group I
category				

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				Prescriber Information	
				Patient Phone:	
rescriber Name: _				Prescriber Phone:	
	NINFORMATION				
NHALED THERAP					
MEDICATION	STRENGTH		D	OSE & DIRECTIONS	QUANTITY/REFILL
Tyvaso (treprostinil) Inhalation Solution	☐ Tyvaso Inhalation System Starter Kit ☐ Tyvaso Refill Kit	breatl breatl			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	Targe 48 per tro Sta daily. every Inh Ot	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:
Yutrepia (Treprostinil) inhalation powder	☐ 26.5 mcg ☐ 53 mcg ☐ 79.5 mcg ☐ 106 mcg	Starting Dosemcg Target Dosemcg Inhale two (2) breaths per capsule, four (4) times daily. Increase by 26.5 mcg, four (4) times daily, every week, as tolerated, to target maintenance dose. Inhale two (2) breaths per capsule,times daily. Increase bymcg,times daily, every week(s)/ day(s) as tolerated, to target maintenance dose.			Quantity: 28-day supply Refills:
NJECTABLE THEF	RAPIES:				
MEDICATION	STRENGTH			DOSE & DIRECTIONS	QUANTITY/REFILLS
☐ Winrevair (sotatercept)	recept) Target Dose (0.7 mg/kg) select one below: Winrevair 45 mg kit (1x45 mg vial) Winrevair 60 mg kit (1x60 mg vial) Winrevair 90 mg kit (2x45 mg vials) Winrevair 120 mg kit (2x60 mg vials) t is interested in patient support programs STAMP SIGNATURE		☐ Inject ml subcutaneously for one dose then increase to ml for target dose after 3 weeks. Dosing interval is every 3 weeks. ☐ Inject ml subcutaneously for dose(s) then increase to ml for target dose after weeks. Dosing interval is every 3 weeks. ☐ Alternative directions;		Quantity: 21-day supply Starter Dose Refills: Target Dose Refills:
	DPRESCRIBER SIGNAT	UKE R	EQUIRED (ST	AMP SIGNATURE NOT ALLOWED)	
"Dispense As Written" / DAW / May Not Substitu Prescriber's Signa			o Substitution /	May Substitute / Product Selection Permitted / Substitution Permissible Prescriber's Signature:	Date:

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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		Patient DOB:	Patient Phone:		
atient Adress:					
rescriber Name: _			Prescriber Phone:		
	NINFORMATION				
IFUSED THERAP		_			
MEDICATION	STRENGTH			NTITY/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	days until goal of	kg/min. Titrate byng/kg/min every ng/kg/min achieved. y days. nps* *For pediatric or low weight patients ONLY over 24 hours kg/min. Titrate byng/kg/min every ng/kg/min achieved. e diluent for Remodulin will be used if no box is n	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	IV infusion continuous Initial dose: ng/days until goal of biluent: Check one (Steril checked) 0.9% NaCl for injection Epoprostenol Sterile description 2 CADD-Legace 2 CVC Care:	over 24 hours kg/min. Titrate byng/kg/min every ng/kg/min achieved. e diluent 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 Initial dose: ng/ days until goal of Discharge dose: ng_ Diluent: Check one (0.9% 0.9% NaCl for injection Pump: 2 CADD-Legac CVC Care: Dressing change ever	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 by ng/kg/min every days until goal of ng/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) □ 0.9% NaCl for injection Sterile Water for injection Pump: □ 2 CADD-Legacy Pumps 2-CADD Solis Pumps CVC Care: Dressing change every days. □ Per IV standard of care			
Patient is interested in pa	<u> </u>	MP SIGNATURE NOT ALLOWED ATURE REQUIRED (ST	Ancillary supplies and kits provided as needed: AMP SIGNATURE NOT ALLOWED)	for administration	
"Dispense As Written" /	Brand Medically Necessary / Do Not S		May Substitute / Product Selection Permitted /		
DAW / May Not Substitu			Substitution Permissible		
Droceriber's Sign	ature:	Date:	Prescriber's Signature:	Date:	

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. 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 communication and any attachments.