## Referral Form for TYVASO (treprostinil) and TYVASO DPI (treprostinil)



Tyvaso and Tyvaso DPI are available only through select Specialty Pharmacy Services (SPS) providers. Follow these 5 steps to complete each section of the following referral form.

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<i>                                      </i>	
(transaction) INNALATION	
ILICUTUSLITIIII SOLUTION	

	CKLIST

- Fill out the Patient and Insurance Information. Let your patient know that an SPS provider will be calling and it is important to answer or return the call.
- □ 2 Complete and sign the Prescriber Information, Medical Information, and Treatment History and Transition Statement.
- □ 3 Complete and sign the Prescription Information, Statement of Medical Necessity for either PH-ILD or PAH, and Calcium Channel Blocker Statement (CCB Statement not required for PH-ILD).
- **4** Complete the Optional Side Effect Management page.
- □ 5 Attach the clinical documents outlined on the Fax Cover Sheet, including right heart catheterization test results, history and physical, and echocardiogram results. Use the included Fax Cover Sheet in this PDF to fax the referral form and signed supporting documents to your preferred SPS provider. (Insurance plans vary and may impact the approval process.)

STEP 1 PATIENT INFORMA	ATION	
Name - First	Middle	Last
Date of Birth	Gender	Last 4 Digits of SSN
Home Address		
City	State	Zip
Shipping Address (if different from home a	nddress)	
City	State	Zip
Telephone: ☐ Home ☐ Cell ☐ Work	Alternate Telephone:   Home   Cell   Work	Best Time to Call: ☐ Morning ☐ Afternoon ☐ Evening Okay to leave a voicemail? ☐ Yes ☐ No
E-mail Address		
Caregiver/Family Member	Caregiver Telephone:   Home   Cell   Work	Caregiver Alternate Telephone:  Home Cell Work
Caregiver E-mail Address	Caregiver Alternate E-mail Address	Okay to leave a voicemail?   Yes No
STEP 1 INSURANCE INFO	RMATION	
Primary Prescription Insurance		
Subscriber ID #	Group #	Telephone

Primary Medical Insurance Policy Holder/Relationship Subscriber ID # Group # Telephone Secondary Medical Insurance Policy Holder/Relationship Subscriber ID # Group # Telephone

Please include copies of the front and back of the patient's medical and prescription insurance card(s).



			Date	of Birth:		
STEP 2 PRESCRIBER INFORM	ATION					
rescriber Name - First Las	st		NPI #		State License #	
Office/Clinic/Institution Name			Office Contac	t Name		
ddress			City		State Zip	
					State Zip	
Office Contact Phone Fax	(		Office Contac	t E-mail		
referred Method of Communication: $\Box$ I	Phone 🔲 E-mail	☐ Mail ☐ Fax				
TEP 2 MEDICAL INFORMATION	ON / PATIENT	EVALUATION	I / SUPPORTING	G DOCUMENTAT	ΓΙΟΝ	
Patient Product Therapy Status for the I	Requested Drug	: Current S	pecialty Pharmacy: Health Group, Inc.	1	Patient Status:  ☐ Outpatient ☐ Inpatient	WHO Grou
NYHA Functional Class (PAH Only):	Weight:ft		Diabetic:  Yes No	Allergies:  Drug Allergies	■ Non-Drug Allergies ■ No	Known Allergie
STEP 2 TREATMENT HISTOR	RY AND TRAN	ISITION STAT	<b>TEMENT</b>			
			Transition State	amont (not required	for DH-TI D. patients)	
Please Indicate Treatment History	Current	Discontinued		ement (not required this patient (if applied		
Please Indicate Treatment History  Medication	Current	Discontinued	It is necessary for <b>FROM</b>	this patient (if appli	icable) to transition	
Please Indicate Treatment History  Medication  PDE-5 I (specify drug(s)):	Current	Discontinued	It is necessary for <b>FROM</b>	this patient (if appl	icable) to transition	
Please Indicate Treatment History  Medication  PDE-5 I (specify drug(s)):  Epoprostenol	Current	Discontinued	It is necessary for <b>FROM</b>	this patient (if appli	icable) to transition	
Please Indicate Treatment History  Medication  PDE-5 I (specify drug(s)):  Epoprostenol  Flolan® (epoprostenol sodium) for Injection	Current	Discontinued	It is necessary for <b>FROM</b>	this patient (if appli	icable) to transition	
Please Indicate Treatment History  Medication  PDE-5 I (specify drug(s)):  Epoprostenol  Flolan® (epoprostenol sodium) for Injection  Letairis® (ambrisentan) Tablets	Current	Discontinued	It is necessary for <b>FROM</b>	this patient (if appli	icable) to transition	
Please Indicate Treatment History  Medication  PDE-5 I (specify drug(s)):  Epoprostenol  Flolan® (epoprostenol sodium) for Injection  Letairis® (ambrisentan) Tablets  Remodulin® (treprostinil) Injection	Current	Discontinued	It is necessary for <b>FROM</b>	this patient (if appli	icable) to transition	
Please Indicate Treatment History  Medication  PDE-5 I (specify drug(s)):  Epoprostenol  Flolan® (epoprostenol sodium) for Injection  Letairis® (ambrisentan) Tablets  Remodulin® (treprostinil) Injection  Tracleer® (bosentan) Tablets	Current	Discontinued	It is necessary for <b>FROM</b>	this patient (if appli	icable) to transition	
Please Indicate Treatment History  Medication  PDE-5 I (specify drug(s)):  Epoprostenol  Flolan® (epoprostenol sodium) for Injection  Letairis® (ambrisentan) Tablets  Remodulin® (treprostinil) Injection  Tracleer® (bosentan) Tablets  Tyvaso® (treprostinil) Inhalation Solution	Current	Discontinued	It is necessary for <b>FROM</b>	this patient (if appli	icable) to transition	
Please Indicate Treatment History  Medication  PDE-5 I (specify drug(s)):  Epoprostenol  Flolan® (epoprostenol sodium) for Injection  Letairis® (ambrisentan) Tablets  Remodulin® (treprostinil) Injection  Tracleer® (bosentan) Tablets  Tyvaso® (treprostinil) Inhalation Solution  Veletri® (epoprostenol) for Injection	Current	Discontinued	It is necessary for <b>FROM</b>	this patient (if appli	icable) to transition	
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Please Indicate Treatment History  Medication  PDE-5 I (specify drug(s)):  Epoprostenol  Flolan® (epoprostenol sodium) for Injection  Letairis® (ambrisentan) Tablets  Remodulin® (treprostinil) Injection  Tracleer® (bosentan) Tablets  Tyvaso® (treprostinil) Inhalation Solution  Veletri® (epoprostenol) for Injection  Ventavis® (iloprost) Inhalation Solution  Adempas® (riociguat) Tablets  Opsumit® (macitentan) Tablets  Orenitram® (treprostinil) Extended-Release  Uptravi® (selexipag) Tablets	Current	Discontinued	It is necessary for <b>FROM</b>	this patient (if appli	icable) to transition	
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Please Indicate Treatment History  Medication  PDE-5 I (specify drug(s)):  Epoprostenol  Flolan® (epoprostenol sodium) for Injection  Letairis® (ambrisentan) Tablets  Remodulin® (treprostinil) Injection  Tracleer® (bosentan) Tablets  Tyvaso® (treprostinil) Inhalation Solution  Veletri® (epoprostenol) for Injection  Ventavis® (iloprost) Inhalation Solution  Adempas® (riociguat) Tablets  Opsumit® (macitentan) Tablets  Orenitram® (treprostinil) Extended-Release  Uptravi® (selexipag) Tablets  Ofev® (nintedanib) Capsules  Esbriet® (pirfenidone) Tablets  Other:		Discontinued	It is necessary for <b>FROM</b>	this patient (if appli	icable) to transition	
Please Indicate Treatment History  Medication  PDE-5 I (specify drug(s)):  Epoprostenol  Flolan® (epoprostenol sodium) for Injection  Letairis® (ambrisentan) Tablets  Remodulin® (treprostinil) Injection  Tracleer® (bosentan) Tablets  Tyvaso® (treprostinil) Inhalation Solution  Veletri® (epoprostenol) for Injection  Ventavis® (iloprost) Inhalation Solution  Adempas® (riociguat) Tablets  Opsumit® (macitentan) Tablets  Orenitram® (treprostinil) Extended-Release  Uptravi® (selexipag) Tablets  Ofev® (nintedanib) Capsules  Esbriet® (pirfenidone) Tablets		Discontinued	It is necessary for <b>FROM</b>	this patient (if appliance)	icable) to transition	
Please Indicate Treatment History  Medication  PDE-5 I (specify drug(s)):  Epoprostenol  Flolan® (epoprostenol sodium) for Injection  Letairis® (ambrisentan) Tablets  Remodulin® (treprostinil) Injection  Tracleer® (bosentan) Tablets  Tyvaso® (treprostinil) Inhalation Solution  Veletri® (epoprostenol) for Injection  Ventavis® (iloprost) Inhalation Solution  Adempas® (riociguat) Tablets  Opsumit® (macitentan) Tablets  Orenitram® (treprostinil) Extended-Release  Uptravi® (selexipag) Tablets  Ofev® (nintedanib) Capsules  Esbriet® (pirfenidone) Tablets  Other:		Discontinued	It is necessary for <b>FROM</b>	this patient (if appliance)	icable) to transition	

Tyvaso and Tyvaso DPI are registered trademarks of United Therapeutics Corporation. All other brands are trademarks of their respective owners. The makers of these brands are not affiliated with and do not endorse United Therapeutics or its products. Please note: The responsibility to determine coverage and reimbursement parameters, and appropriate coding for particular patient and/or procedure, is the responsibility of the provider. The information provided here is not a guarantee of coverage or reimbursement.

Patient Na	me: Date of Birth:	
STEP 3	PH-ILD - USE THIS SECTION FOR PH-ILD	
Diagnosis -	The following ICD-10 codes do not suggest approval, coverage, or reimbursement for specific uses or ind	lications.
PH Diagnosi ICD-10 I27.23 ILD Diagnos IIP: ICD-10 CTD-related Environmen ICD-10 J61	Pulmonary hypertension due to lung diseases and hypoxia Other ICD-10:	112 Idiopathic pulmonary fibrosis
	Please visit www.utassist.com/codes for additional ICD-10 codes related to PAH, PH, and	ILD
	TYVASO (treprostinil) 1.74mg/2.9ml ampule (0.6mg/ml) Inhalation Solution  Target dose: 9 breaths (54 mcg) to 12 breaths (72 mcg), 4 times daily - Start with 3 breaths (18 mcg) 4 times daily (if 3 breaths are not tolerated, use 1 to 2 breaths). Increase by an additional 1 breath per week, as tolerated, until the target dose of 9 breaths (54 mcg) to 12 breaths (72 mcg), 4 times daily is achieved.  TYVASO Inhalation System Starter Kit (28-day supply) 0 refills TYVASO Inhalation System Refill Kit (28-day supply) X refills  Prescriber may specify any alternative or additional dosing and titration instructions here:	Dose Comparison  TYVASO Nebulizer # of Breaths  ≤5  TYVASO DPI Cartridge Strength
	TYVASO DPI (treprostinil) Inhalation Powder  Target dose: 48 mcg or 64 mcg or Other mcg per treatment session, 4 times daily (Check One)  Start with one 16-mcg cartridge per treatment session, 4 times daily. Increase cartridge strength by 16 mcg per treatment session every 1 to 2 weeks, as tolerated, to selected target dose. Titration schedule may vary based on tolerability. If the prescribed dose is higher than 64 mcg per treatment session, more than 1 cartridge will be needed per session.  TYVASO DPI Titration Kit (28-day supply) Choose for titration phase.  16 mcg (112 ct), 32 mcg (112 ct), and 48 mcg (28 ct) 1 refill  TYVASO DPI Maintenance Kit (28-day supply) X refills	6 to 7 32 mcg 8 to 10 48 mcg 11 to 12 64 mcg
	Inhale one breath per cartridge, 4 times daily. Please check the box of the maintenance kit for the desired target do 16 mcg (112 ct) 32 mcg (112 ct) 48 mcg (112 ct) 64 mcg (112 ct)  Prescriber may specify any alternative or additional dosing and titration instructions on the line below. If the per treatment session, more than 1 cartridge will be needed per session:  Specially Pharmacy to contact prescribing practitioner for adjustments to the written orders specified above.	
specific require  RECK Nurse Visits	ORDERS  RN visit to provide assessment and education on administration, dosing, and titration. Location:  It is to comply with their state-specific prescription requirements, such as e-prescribing, state-specific prescription form, fax languagements could result in outreach to the Prescriber.  OR Prescriber directed Specialty Pharmacy home healthcare RN visit to provide education on self-administration of Tyvaso or Tyvaso DPI, including dose, titration, and side effect management.	uage, etc. Non-compliance of state-
ERE PHYS	PRESCRIBER SIGNATURE: PRESCRIPTION AND STATEMENT OF MEDICAL NECESSITY for that the pulmonary hypertension associated with interstitial lung disease therapy ordered above is medically necessary and that I am SICIAN'S SIGNATURE REQUIRED TO VALIDATE PRESCRIPTIONS.	personally supervising the care of this patient
Physic	cian's Signature: Dispense as Written Substitution Allowed	Date:
AW	Specific Dispense as Written (DAW) Selection Verbiage:	

Patient Na	me: Date of Birth:	
STEP 3	PAH - USE THIS SECTION FOR PAH	
Diagnosis -	The following ICD-10 codes do not suggest approval, coverage, or reimbursement for specific uses or indication	ens.
ICD-10 I27.0	Primary pulmonary hypertension: 🗌 Idiopathic PAH 🗎 Heritable PAH	
ICD-10 I27.21	L Secondary pulmonary arterial hypertension:  Connective tissue disease Drugs/Toxins induced Portal hypertension	HIV Congenital heart diseases
Other:	Other ICD-10:	
	Please visit www.utassist.com/codes for additional ICD-10 codes related to PAH, PH, and ILD	
	TYVASO (treprostinil) 1.74mg/2.9ml ampule (0.6mg/ml) Inhalation Solution  Target dose: 9 breaths (54 mcg) to 12 breaths (72 mcg), 4 times daily - Start with 3 breaths (18 mcg) 4 times daily (if 3 breaths are not tolerated, use 1 to 2 breaths). Increase by an additional 3 breaths every week, if tolerated, until the target dose of 9 breaths (54 mcg) to 12 breaths (72 mcg), 4 times daily.  TYVASO Inhalation System Starter Kit (28-day supply) 0 refills  TYVASO Inhalation System Refill Kit (28-day supply) X refills  Prescriber may specify any alternative or additional dosing and titration instructions here:	TYVASO Nebulizer # of Breaths Strength
	OR	≤5 16 mcg
	TYVASO DPI (treprostinil) Inhalation Powder	6 to 7 32 mcg
	Target dose: 48 mcg or 64 mcg or 6ther mcg per treatment session, 4 times daily (Check One)  Start with one 16-mcg cartridge per treatment session, 4 times daily. Increase cartridge strength by 16 mcg per treatment	8 to 10 48 mcg 📙
	session every week to selected target dose. Titration schedule may vary based on tolerability. If the prescribed dose is higher than 64 mcg per treatment session, more than 1 cartridge will be needed per session.	11 to 12 64 mcg
	TYVASO DPI Titration Kit (28-day supply) Choose for titration phase.	
	16 mcg (112 ct), 32 mcg (112 ct), and 48 mcg (28 ct) 1 refill	
	TYVASO DPI Maintenance Kit (28-day supply) X refills  Inhale one breath per cartridge, 4 times daily. Please check the box of the maintenance kit for the desired target dose.	
	☐ 16 mcg (112 ct) ☐ 32 mcg (112 ct) ☐ 48 mcg (112 ct) ☐ 64 mcg (112 ct)	
	Prescriber may specify any alternative or additional dosing and titration instructions on the line below. If the prescri per treatment session, more than 1 cartridge will be needed per session:	bed dose is higher than 64 mc
	Specialty Pharmacy to contact prescribing practitioner for adjustments to the written orders specified above.	
specific require THECK Nurse ONE Visits	RN visit to provide assessment and education on administration, dosing, and titration. Location: Home is to comply with their state-specific prescription requirements, such as e-prescribing, state-specific prescription form, fax language, elements could result in outreach to the Prescriber.  Specialty Pharmacy home healthcare RN visit to provide education on self-administration of Tyvaso or Tyvaso DPI, including dose, titration, and side effect management.	tc. Non-compliance of state-
STEP 3	CALCTUM CHANNEL BLOCKER STATEMENT (Not required for BU TLD notionts)	
	<i>,</i>	o voculto
	te below if the Patient named above was trialed on a Calcium Channel Blocker prior to the initiation of therapy and indicate the Channel Blocker was not trialed because:	e resuits.
	as depressed cardiac output Patient is hemodynamically unstable or has a history of postural hypotension	
	as systemic hypotension Patient did not meet ACCP Guidelines for Vasodilator Response	
	as known hypersensitivity Patient has documented bradycardia or second- or third-degree heart block	
OR		
The followi	ng Calcium Channel Blocker was trialed:	
	owing response(s):	
☐ Patient hy ☐ Adverse €	/persensitive or allergic Pulmonary arterial pressure continued to rise event Patient became hemodynamically unstable	
	ontinued to progress or patient remained symptomatic	
Other: _		
STEP 3	PRESCRIBER SIGNATURE: PRESCRIPTION AND STATEMENT OF MEDICAL NECESSITY	
I certi	fy that the pulmonary arterial hypertension therapy ordered above is medically necessary and that I am personally supervising the care	e of this patient.
HERE	SICIAN'S SIGNATURE REQUIRED TO VALIDATE PRESCRIPTIONS.	D-4
Physic	cian's Signature: Dispense as Written Substitution Allowed	Date:
DAW State-	Specific Dispense as Written (DAW) Selection Verbiage:	

(Physician attests this is his/her legal signature. NO STAMPS.) PRESCRIPTIONS MUST BE FAXED.

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tient Name:	Date of Birth:
STEP 4 OPTIONAL SIDE EFF	ECT MANAGEMENT
By providing your side effect management str dosing in Step 3 of this form.	rategies, SPS will be able to follow up with the patient should they experience side effects. Include directions to SPS for
*INFORMATION PROVIDED BELOW IS NOT A PRES	SCRIPTION; RATHER, IF ADDITIONAL PRESCRIPTIONS ARE INTENDED, THEY SHOULD BE PROVIDED TO THE PATIENT SEPARATELY.
Headache:	
Acetaminophen mg	Frequency Opioids (separate Rx required) Tramadol (separate Rx required)
■ NSAIDs (separate Rx may be required	d)
Other	
Nausea/Vomiting:	
Ondansetron (separate Rx required)	☐ Metoclopramide (separate Rx required) ☐ PPIs (separate Rx may be required)
Prochlorperazine (separate Rx required	d) Promethazine (separate Rx required)
Remind patient to hold the device level ar	nd swish & spit after each treatment session
Other	
Throat Irritation:	
$\square$ Oral phenol-based analgesic sprays $\square$ Re	eview medication administration technique
☐ Other	
Cough:	
Albuterol (separate Rx required) Be	enzonatate (separate Rx required) — Cough suppressant (separate Rx may be required)
	ozenges (note: not to be used during treatment session)   Inhaled anticholinergics (separate Rx required)
■ Inhaled steroids (separate Rx required)	Other
Diarrhea:	
$\square$ Loperamide (separate Rx required) $\square$	Other
Address of war are set as	
Additional Instructions:  Provide any additional instructions for SPS of	on preferred communication or managing other side effects.
Frovide any additional instructions for SF3 of	in preferred communication of managing other side effects.
	_

Fax the completed referral form and documentation to the Specialty Pharmacy of your choice below.

Date:	Patient Initials:	Patient Date of Birth:
To: (check one)	☐ <b>Accredo Health Group, Inc.</b> Fax: 1-800-711-3526 Phone: 1-866-344-4874	□ <b>CVS Specialty</b> Fax: 1-877-943-1000 Phone: 1-877-242-2738
From: (Name of agent o	f prescriber who transmitted the facsimile/prescription	n)
Facility Name:		
Fax:		
Included in this fa	Y'	
	nso and Tyvaso DPI Therapy Referra	al Form including
<ul><li>Step 1 - Patient 1</li><li>Step 2 - Prescrib</li></ul>	Information and Insurance Information (including from er Information, Medical Information/Patient Evaluatio tion Information and Calcium Channel Blocker Statem	nt and back copies of medical and prescription insurance card(s)) n/Supporting Documentation, and Treatment History and Transition Statement
<ul> <li>Step 4 - Optiona</li> </ul>	i Side Effect Management	
	l and dated documents	
<ul> <li>Included signed</li> <li>Right Heart Cath</li> <li>History and Phys</li> <li>Need for Specific</li> <li>Echocardiogram</li> </ul>	l and dated documents eterization Results	ated with ILD Clinical Signs and Symptoms, Course of Illness) ute walk test not required for PH-ILD)
<ul> <li>Right Heart Cath</li> <li>History and Phys</li> <li>Need for Specific</li> <li>Echocardiogram</li> <li>High-Resolution</li> </ul>	l and dated documents eterization Results cical (including Onset of Symptoms, PAH or PH associate Drug Therapy and 6-minute walk test results (6-minute Results (not required for PH-ILD patients)	ute walk test not required for PH-ILD)
<ul> <li>Right Heart Cath</li> <li>History and Phys</li> <li>Need for Specific</li> <li>Echocardiogram</li> <li>High-Resolution</li> </ul> Number of Pages:	l and dated documents eterization Results sical (including Onset of Symptoms, PAH or PH associa: Drug Therapy and 6-minute walk test results (6-min Results (not required for PH-ILD patients) CT Scan (not required for PAH patients)	ute walk test not required for PH-ILD)
<ul> <li>Right Heart Cath</li> <li>History and Phys</li> <li>Need for Specific</li> <li>Echocardiogram</li> <li>High-Resolution</li> </ul>	l and dated documents eterization Results sical (including Onset of Symptoms, PAH or PH associa: Drug Therapy and 6-minute walk test results (6-min Results (not required for PH-ILD patients) CT Scan (not required for PAH patients)	ute walk test not required for PH-ILD)

**US-TYV-0710** 

