# Practical considerations for managing transitions from parenteral prostacyclins to an oral IP receptor agonist

Jean M Elwing, MD¹; Christina Benninger, MSN, APRN²; Paul Strachan, MD²; Korawin Triyasakorn, PharmD²; Marla Choren, PharmD, MPH²; Kody Timms, PharmD²; Lori Reed, NP-C³

<sup>1</sup>University of Cincinnati, Cincinnati, OH, USA; <sup>2</sup>Johnson & Johnson, Titusville, NJ, USA; <sup>3</sup>Piedmont Healthcare, Atlanta, GA, USA

#### Introduction

- The prostacyclin pathway represents a key foundational pathway in the treatment of patients with pulmonary arterial hypertension (PAH), with over 30 years of clinical experience—including 10 years with an oral IP receptor agonist<sup>1–3</sup>
- Parenteral prostacyclin pathway agents (PPAs) are often used for the treatment of patients at high risk, or for those who remain at intermediate/high risk on initial therapy<sup>4</sup>
- Parenteral PPA use may be limited by administration difficulties, including the need for continuous infusion, administration side effects, and complexities of a chronic indwelling catheter
- An alternative route of therapy, such as oral administration, may be a more favorable option for some patients

# • Methods for transitioning to an oral IP receptor agonist remain undefined as treatment protocols are lacking

 A systematic literature review revealed a high level of heterogeneity in approaches to transitioning from a parenteral PPA to an oral IP receptor agonist (Elwing et al, PH Hope 2025 Poster), highlighting a need for standardized, evidence-based, practical recommendations for managing this transition

# Objective

 To describe practical considerations for managing patients who are transitioning from a parenteral PPA to an oral IP receptor agonist based on the expertise of two healthcare providers

## Results

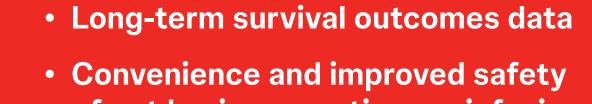
### SELECT PPA BASED ON PROS AND CONS

#### **PARENTERAL**



- Long-term outcomes data from registries
- Rapid dose-adjustment effects with no maximum dose
- Implied adherence because of administration route
- Complexities and challenges
   associated with continuous infusion and
   equipment, including line infections, loss
   of lines, line maintenance, and
   subcutaneous catheter site pain
- Need for additional care partner, social, and nursing support; close monitoring; and management of the pump and lines, and systemic adverse effects
- Medication side-effects, uptitration, and administration
- Limited long-term care or rehab options if facilities are not familiar with the medication required for inpatient management
- Education required for care facility staff and care partners

#### ORAL IP RECEPTOR AGONIST



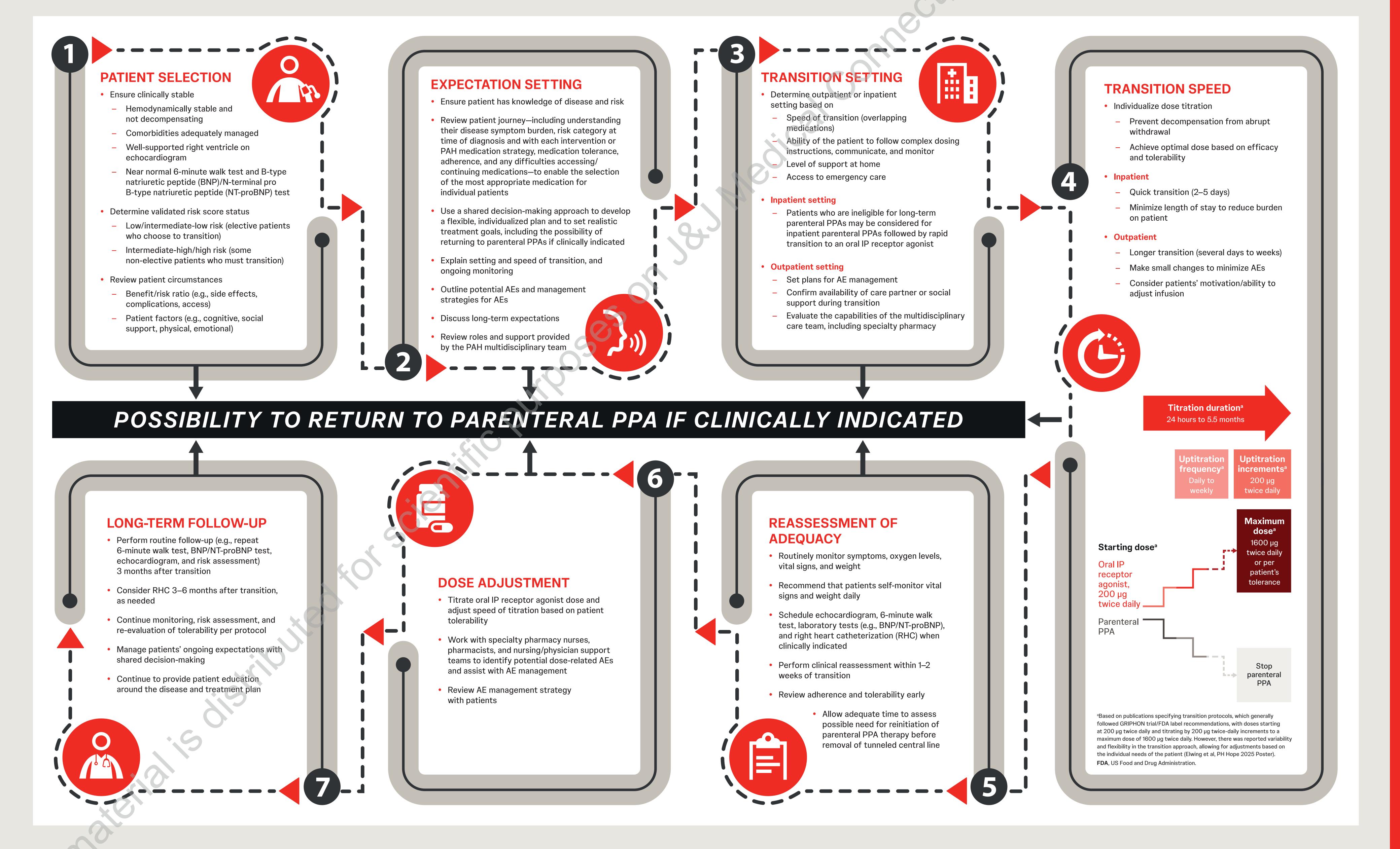


- of not having a continuous infusion or central line, pumps, or catheters (intravenous or subcutaneous)
- Fewer safety concerns if medication is abruptly discontinued
- Well-established dosing with a personalized goal dose
- Manageable side effect profile
- Potential for improved quality of life



- Frequent management and monitoring with dose titration
- Potential side effects and adverse events (AEs)
- Possible decreased adherence with intermittent, twice-daily dosing schedule

# CONSIDERATIONS FOR TRANSITIONING FROM PARENTERAL PPA TO ORAL IP RECEPTOR AGONIST



# Key takeaway

Parenteral PPA to an oral IP receptor agonist can be a challenging and complex process.
However, a patient's opportunity for a successful transition is maximized through careful planning, setting transparent expectations, and an individualized approach with close monitoring and long-term follow-up

# Conclusions

- Careful selection of patients is imperative to successfully transition from a parenteral PPA to an oral IP receptor agonist
- It is important to set transparent and realistic expectations through an individualized, shared decision-making approach with patients and involvement of a multidisciplinary team
- Patients should be informed of the possibility of returning to parenteral PPAs throughout the process, if clinically indicated
- Close monitoring during the transition, immediately after completion, and long-term follow-up are essential for successful transition

#### Acknowledgments

This study was funded by Johnson & Johnson. Medical writing support was provided by Ashley Sizer, PhD, CMPP, on behalf of Twist Medical and was funded by Johnson & Johnson.

#### Disclosures

JME has served as a consultant for United Therapeutics, Aerovate, Gossamer Bio, Liquidia, Merck, Johnson & Johnson, Insmed, and Pulmovant. She has also participated in clinical research studies funded by United Therapeutics, Gossamer Bio, Bayer, Acceleron/Merck, Altavant, Aerovate, Pharmosa/Liquidia, Johnson & Johnson, Lung LLC, and Pulmovant. CB, PS, K Triyasakorn, MC, and K Timms are employees of Johnson & Johnson. LR has served as a consultant for Johnson & Johnson and United Therapeutics. She has also participated in speaker bureaus for Johnson & Johnson, Liquidia, Merck, and United Therapeutics.

Pulmonary Hypertension





Scan the QR code

The QR code is intended to provide scientific information for individual reference, and the information should not be altered or reproduced in