

# The PDP Reach: How Far Should PDPs Extend?

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# Current Role of Access within PDPs

- **Access imbedded in work of organization**
  - R&D, BD, Executive follow access principles
  - Focus of the Access Teams
- **Access informs product development**
  - How access considerations are incorporated into development of new product
  - How access informs target product profiles (TPP)
  - Consider: Mechanisms for mapping user requirements & Timing – before Phase 3, know the market
- **Access facilitates timely uptake**
  - Think through all the needs and facilitate potential roadblocks: fewer surprises
  - Catalyze/Leverage other partners
  - Fill gaps



# PDP Role in Post-Licensure

- **Prioritize activities at country level**
  - Financing - the biggest hurdle?
  - Regulatory
  - Policy change
  - Pricing
  - Distribution/sales
  - Patient/provider education
- **Define Objectives: How far should we go?**

SRA  
regulatory  
approval

WHOPQ

WHO  
“endorsement”  
of the product

Regulatory or  
policy approval  
at country level

Product  
purchase

The last mile



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# Partners

- **Are the partners in place to take the rest on?**
  - Partners with offices: e.g. MVI has PATH, others have...?
  - Manufacturers: relationships with regulatory, distribution, sometimes govt relations
  - Technical assistance providers in disease areas, supply chain management
  - WHO
  - National programs
- **Are these sufficient or are there gaps?**
- **Input from countries and WHO needed!**



# Defining Our Mandate:

## What is appropriate PDP Role Post-Licensure?

- **Do we have the expertise?**
  - Better to highlight the relevant gaps and advocate for others to take action?
- **What successful balancing is currently happening?**
  - Lessons learned from those on the market now
- **How can we be clearer about what we are doing?**
  - Is any of us covering the last mile?
  - If the worry is about cost, perhaps we need to define what costs we expect to have at implementation stage
  - Input from donors needed!

