

## MPT Priority Action Areas & Gaps

As of September 2022, the priority **MPT Fieldwide Action Areas** identified by the SACC are as follows:

### 1. **A productive ecosystem of MPT product R&D**

- Ecosystem expansion - Part 1: Expanding MPT R&D with a focus on new products
  - Expanded pipeline of researchers for expanded product R&D
  - Enhanced physical infrastructure for product R&D
  - Geographic expansion for MPT R&D in terms of primary researchers and collaborative efforts
  - Expanded generic licensing opportunities and geographically broad manufacturing base
- Ecosystem expansion - Part 2: Achieving appropriate funding to support necessary expansion of MPT ecosystem
  - Novel and innovative funding and investment approaches to ensure successful development of MPTs through commercialization and scale-up
  - Enhanced ability for cost-effective collaborations

### 2. **Improved understanding of reproductive biology for the purpose of new pharmaceutical development for MPT R&D**

- Enhanced focus on non-hormonal MPTs
- Enhanced focus on male contraceptive options
- Expanded understanding of cervicovaginal microbiome, pharmacogenomics, etc.
- Expanded understanding of underlying conditions in those likely to use MPTs (e.g., perimenopause, cognitive disabilities, other underlying health issues)

### 3. **Expanded understanding of socio-behavioral research (SBR) considerations among underrepresented groups in MPT research, including**

- People under the age of 18
- People with gynecological conditions
- Sexual and gender minority (SGM) populations
- Older populations (e.g., perimenopausal women)
- Those known to metabolize drugs differently (e.g., obese individuals, Down syndrome individuals)

### 4. **Expanded understanding of market considerations to help ensure successful commercialization and uptake of MPTs, including**

- Market segmentation and target segment identification
- Switching between products and triggers estimates
- Demand forecasting

### 5. **Enhanced understanding of innovative approaches for MPT clinical trials that address regulatory and ethical challenges of testing multiple indications in the same trial**