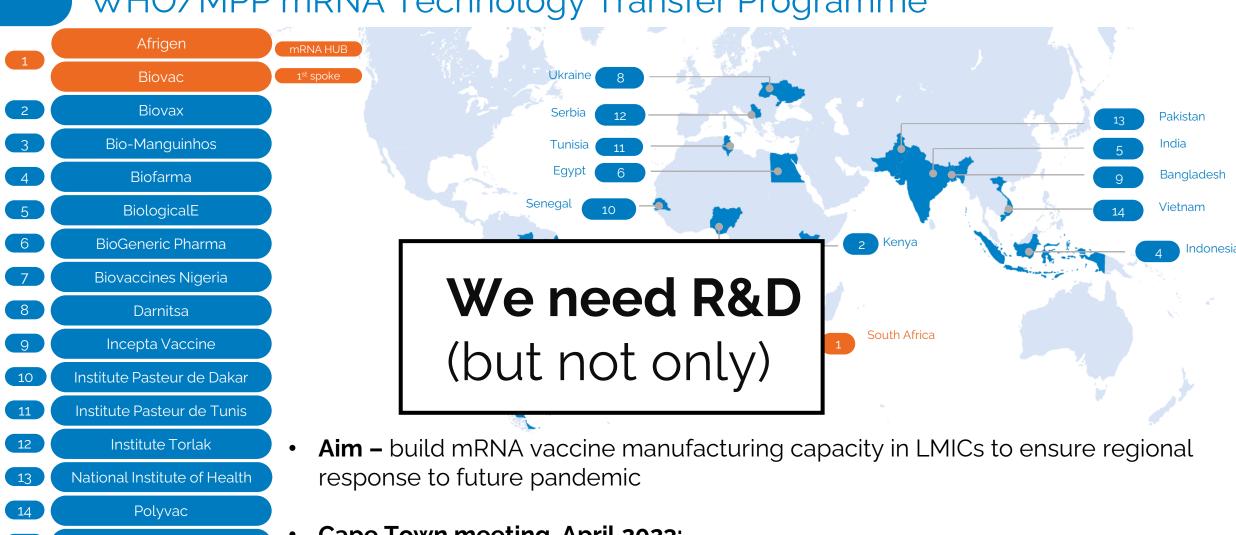
WHO/MPP mRNA Technology Transfer Programme



- Cape Town meeting, April 2023:
 - Establishing mRNA manufacturing capacity in not sufficient In absence of routine manufacturing activity these facilities will not be sustainable

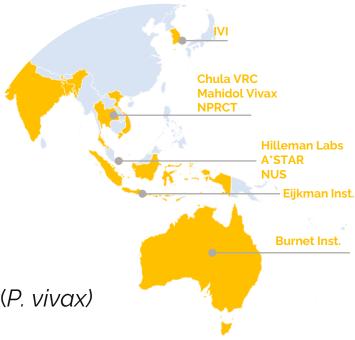


Sinergium Biotech

Towards establishing R&D consortia

- R&D regional collaborative approach
 - Regional PH problem/market and set of key stakeholders
- Identify mRNA products that are commercially viable
 - Based on country needs
 - PTRS/PPDP evaluation
- Identify R&D centers to drive mRNA product development
 - Identify key R&D collaborators (e.g. academics, translational)
 - Recognizing that manufacturers have limited R&D capacity
- Bangkok scientific consultation, October 2023
 - Dengue, HFMD, Tx HPV, P. vivax
 - Establish consortia to drive mRNA product development
 - IVI (dengue), Hilleman (HFMD), Chula VRC (HPV), Mahidol (P. vivax)
- Replicate the model in other WHO regions







R&D consortia – High-level Considerations

- Consortium Goal develop commercially viable mRNA products that can be manufactured by the WHO/MPP mRNA manufacturing partners
 - Evidence-based product development up to proof-of-concept early clinical trial and transfer to manufacturing partners for further development
 - Locally owned
 - Consortia leads identify partners and responsible for resource mobilization
 - Not WHO/MPP TT programme, but WHO/MPP to support as needed
- Ensure ACCESS to all WHO/MPP partners.
 - Technology should be made available to all partners
 - Reasonable royalties not a barrier.
 - Open to innovations
 - Compatibility with Afrigen process
 - Sharing/access to technology for candidate vaccines should permit recipients to adapt to their manufacturing process/technology
 - Singapore Declaration of Commitment



