Full TechWatch Presentation Guide (1 hour)

Please review the information below before creating a presentation for your <u>Full TechWatch</u> meeting. The information will guide you, so you can have a productive and interactive meeting with BARDA's staff. During your meeting, BARDA staff will engage in technical discussion during your presentation to provide guidance, seek clarification, or ask questions.

Timeline:

- One week prior to your meeting: send your draft presentation to the TechWatch team to review.
 - Please refer to your Meeting Invitation email for TechWatch Team contact info.
- At least 24-48 hours prior to your meeting:
 - Send your final presentation as a PowerPoint or PDF file to the TechWatch Team to allow USG colleagues time to review your slides in advance, AND
 - Upload your final presentation materials to the <u>BDR</u> Stakeholder Portal.
- On the day of the meeting: please arrive 5 minute early to the meeting for the initial meeting frame-in, and to ensure an on time start.

General Guidance for Full TechWatch:

- The meeting will be 60 minutes in duration; 45 minutes max for presentation (includes technical discussion throughout) and 15 minutes for Q&A and closeout discussion.
- The presentation should be data-driven and should not exceed 30-40 slides
- Explain how your technology aligns with the requirements stated in a specific area of interest (AOI) in BARDA's <u>BAA</u> or <u>EZ-BAA</u>.
- Review information on <u>Technology Readiness Level</u> to determine the stage of maturity of your development program.

Specific Slide Guidance for Full TechWatch:

Slide Content	# of Slides
Company Overview	1
Key Personnel	1
Product Pipeline Overview	1
Alignment	1
Specific AOI and/or BARDA's BAA or EZ-BAA, and TRL	1
Animation	1
If possible, of how your technology works (i.e. MOA)	1
Proof-of-Concept Data	8-12
Product Differentiation	
Comparison with other competing technologies to define value proposition/	1
ROI to BARDA/USG (i.e., table where rows are attributes, columns are your	1
technology vs. others; 3-4 max)	
Regulatory	
FDA feedback on development plan, and regulatory strategy for	1-2
approval/licensure/clearance	
Clinical Data/Development Plan	6-8
Manufacturing	1-2
Intellectual Property	1
Proposed Business Plan	1
For commercial sustainability if applicable	1
Funding Landscape	
High level review of estimated funding required to complete the development	
program/establish the capability through FDA licensure/approval/clearance	
Key Questions for BARDA	1
3-4 key questions	1
Contact Details	1
Name, phone number, email address, website	1