Patient Experience Data in Regulatory Processes:

Informing stakeholder-regulator communication toward medical product development

June 25th – 15h30 CET / 9.30am EST

Objectives:

- Learn more about ongoing patient engagement efforts to inform regulatory decision making
- Explore challenges and emerging good practices around the use of patient experience data in regulatory submissions
- Prioritize Patient Experience Data (PED) challenges and emerging good practices in regulatory communication
- Understand what areas can help move the needle internally for organizations and externally in collaboration with patient organizations, biopharma and regulators