



## **Patient Experience Data in Regulatory Processes:**

**Informing stakeholder-regulator communication toward medical product development**

*June 25<sup>th</sup> – 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