

Participant & Community Engagement – Potential Key Performance Indicators (KPIs)

Output indicators

- Process established for target subpopulation(s) to voice inclusion during trial design
- Number of partnerships established with patient advocacy and community organizations relevant to target subpopulation(s)
- Number of in-person meetings held with patients of target subpopulation(s) to guide study design and recruitment planning
- Proportion of participant-facing materials reviewed by patients and advocates of target subpopulation(s)
- Proportion of clinical trials with patient feedback processes at the end-of-study with patients of target subpopulation(s)
- Community advisory boards established for target subpopulation(s)

Outcome indicators

- Adjustment to study protocol and/or recruitment materials made based on target subpopulation(s) engagement activities
- Target subpopulation(s) input represented at annual review
- Proportion of subpopulation(s) relationship sustained across more than one trial