**Patient & Community Engagement in the Design and Implementation of Research Studies Resource Guide**

**Tracking Form**

**Level 2: Apply**

**Name:** Click or tap here to enter text.

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| **Order** | **Objectives** | **Status** | **Comments** |
| **1** | Meaningfully **supporting patient and community partners in research** and related processes | [ ]  Started[ ]  Completed[ ]  Plan to start in the future[ ]  Not applicable |  |
| **2** | Best practices for including **people with lived experience** in research teams  | [ ]  Started[ ]  Completed[ ]  Plan to start in the future[ ]  Not applicable |  |
| **3** | Understanding the **concept of power sharing** (role of power and self-location in community engagement) | [ ]  Started[ ]  Completed[ ]  Plan to start in the future[ ]  Not applicable |  |
| **4** | Sharing **budgets and resources** with patient and community partners  | [ ]  Started[ ]  Completed[ ]  Plan to start in the future[ ]  Not applicable |  |
| **5** | Ensuring a **collaborative approach to knowledge mobilization** between patient/community and researcher  | [ ]  Started[ ]  Completed[ ]  Plan to start in the future[ ]  Not applicable  |  |
| **6** | Best practices for **creating equitable research teams** when engaging patients and community partners  | [ ]  Started[ ]  Completed[ ]  Plan to start in the future[ ]  Not applicable  |  |
| **7** | Ensuring an **anti-racist, equitable and socially accountable approach** to recruitment of community members, patients, family, and people with lived experience | [ ]  Started[ ]  Completed[ ]  Plan to start in the future[ ]  Not applicable  |  |