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| **Screening Date** | | **Screening Attempt** | **PTID** | | | **Staff Initials/Date** | | | **Enrollment Date  (or N/A if**  **not enrolled)** | **Screen Failure Date (or N/A if enrolled)** | | | **Screening Failure/ Discontinuation Codes**  **(or N/A if enrolled)** | | | **Staff Initials/Date** |
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| **Screening Failure/Discontinuation Codes**  ϕ Individuals who can get pregnant | | | | | | | | | | | | | | | | |
| **I1** | Not cis- or transgender man or TGW aged 18-35 (inclusive) | | | **I7** | No RAI/inadequate RAI frequency | | **E2** | Use of non-study rectally-administered products/meds | | | **E8** | Condomless RVI/RAI while not on PrEP with partner of unknown HIV status and unknown PrEP/ART use, within 1 month | | **N1** | Other – Declines enrollment | |
| **I2** | Not able/willing to provide IC | | | **I8** | Not willing to refrain from participation in other studies | | **E3** | Known adverse reaction to study product | | | **E9** | IVDU within 12 months | | **N2** | Other – No enrollment w/in 45-day window | |
| **I3** | HIV 1/2 positive | | | **I9** | Positive pregnancy test ϕ | | **E4** | Participation in another study within 30 days of Enrollment | | | **E10** | Diagnosed w/ RTI/STI/UTI | | **N3** | Other: Accrual target met | |
| **I4** | Inadequate locator information | | | **I10** | No contraception ϕ | | **E5** | Participation in rectal product study (ever) | | | **E11** | Pregnant/breastfeeding or intends to become pregnant/breastfeed ϕ | | **N4** | Other: | |
| **I5** | Not able/willing to comply with study requirements | | | **E1a** | History of inflammatory bowel disease (IBD) | | **E6** | PEP within 3 months | | | **E12** | Pregnancy outcome within 90 days ϕ | |  |  | |
| **I6** | Not in general good health | | | **E1b** | Exclusionary anorectal condition | | **E7** | Condomless RVI/RAI while not on PrEP with HIV+ partner who is not on ARV or of unknown ARV status, within 3 months | | | **E13** | Any other condition, per IoR/designee | |  |  | |

ϕ Individuals who can get pregnant