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| **ScreeningDate** | **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** |
| **I-1** | Not between 16-21 yrs old  | **I-8** | Positive pregnancy test | **E-1d** | Plans to travel away | **E-5d** | Injection drug use in previous 12 months  | **E-6b** | Hemoglobin Grade 2 or higher |
| **I-2** | No informed consent/ assent/ parental permission | **I-9** | No effective contraception | **E-2** | HIV-positive test (at least 1 positive rapid) | **E-5e** | Use of HIV PEP and/or PrEP w/in 3 months prior | **E-6c** | Creatinine Clearance < 60 mL/min |
| **I-3** | Inadequate locator | **I-10** | Not willing to refrain from vaginal products/practices 72 hrs prior to visits | **E-3** | Diagnosed with UTI/RTI/ STI  | **E-5f** | Breastfeeding | **E-7** | Any other condition (per IoR/designee) |
| **I-4** | Not willing to comply with study | **I-11** | Not willing to refrain from other studies | **E-4** | Pelvic finding grade 2 or higher | **E-5g** | Pregnancy outcome w/in 8 last weeks | **N-1** | Other – Declines enrollment |
| **I-5** | Pre-menarche | **E-1a** | Plans to get pregnant | **E-5a** | Known study product adverse reaction | **E-5h** | Participation in research study w/in last 60 days | **N-2** | Other – Exceeds 70-day enrollment window |
| **I-6** | HIV infected (per 2 positive rapids or confirmation test) | **E-1b** | Intends to access/use oral PrEP outside study | **E-5b** | Known latex/polyurethane adverse reaction | **E-5i** | Significant uncontrolled active disease (per IoR/designee) | **N-3** | Other: |
| **I-7** | No history of sexual intercourse | **E-1c** | Plans to relocate | **E-5c** | Symptomatic for acute HIV infection | **E-6a** | Positive HBsAG  | **N-4** | Other: |