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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** | | | | | | | | | | | | | | | |
| **I-1** | Not between 16-21 yrs old | | | **I-9** | No effective contraception | | **E-2** | HIV-positive test | **E-5e** | | Use of HIV PEP and/or PrEP w/in 3 months prior | | **E-6c** | Creatinine Clearance < 60 mL/min | |
| **I-2** | No informed consent/ assent/ parental permission | | | **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 (IoR/designee) | |
| **I-3** | Inadequate locator | | | **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-4** | Not willing to comply with study | | | **E-1a** | Plan to get pregnant | | **E-5a** | Known study product adverse reaction | **E-5h** | | Participation in drug/device/vaginal product/vaccine trial w/in last 60 days | | **N-2** | Other – No enrollment visit within 70-day window | |
| **I-5** | Pre-menarche | | | **E-1b** | Intends to access/use oral PrEP outside study | | **E-5b** | Known latex/polyurethane adverse reaction | **E-5i** | | Significant uncontrolled active disease (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: | |
| **I-8** | Positive pregnancy test | | | **E-1d** | Plans to travel away | | **E-5d** | IDU in previous 12 months | **E-6b** | | Hemoglobin Grade 2 or higher | |  |  | |