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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** | | | | | | | | | | | | | | | | |
| **I1** | Not between 18-45 yrs old  Not born female | | | **I8** | Not in good health | | **E1** | BMI greater than 35 kg/m2 | | **E6c** | | Contraindication to progestin | | E6j | Pregnancy outcome w/in 90 days | |
| **I2** | No informed consent | | | **I9** | HIV-positive | | **E2** | Pregnant or intends to be pregnant | | **E6d** | | Use of hormonal contraception | | E6k | Genital procedure w/in 60 days | |
| **I3** | Inadequate locator | | | **I10** | Non regular menstrual cycles | | **E3** | Diagnosed with UTI/RTI | | **E6e** | | Chronic use of prohibited meds | | E6i | Breastfeeding | |
| **I4** | Non-English speaking | | | **I11** | Non intact uterus with at least one ovary | | **E4** | Diagnosed with STI | | **E6f** | | DMPA use in 6 months prior | | E6m | Participation in drug/device trial | |
| **I5** | Not willing to comply with study | | | **I12** | Not willing to abstain from vaginal products | | **E5** | Pelvic finding grade 2 or higher | | **E6g** | | IDU in 12 months prior | | E7 | Grade 1 or higher abnormal lab | |
| **I6** | Not willing to abstain from sex | | | **I13** | ≥21 y.o: unsatisfactory Pap | | **E6a** | Known adverse reaction to products | | **E6h** | | Use of PEP | | E8 | Any other condition (IoR/designee) | |
| **I7** | Non-effective contraception | | | **I14** | Not willing to refrain from other studies | | **E6b** | Chronic vaginal candidiasis | | **E6i** | | Use of PrEP | |  |  | |