| **Visit 17 (Follow-Up Safety Contact and Termination Visit)** |
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| **Procedures** | **Staff****Initials** |
|  | Confirm identity and PTID. |  |
|  | Review/update locator information per site SOPs. |  |
|  | Document any adverse events: Complete/update AE Log CRF(s) as needed. If indicated, schedule interim visit for follow-up of identified AEs. |  |
|  | Provide and explain available exam and lab test results. If STI/RTI/UTI is diagnosed, provide or refer for treatment. Document provision of results, treatments and referrals in chart notes. |  |
|  | Reinforce site contact information and: * If applicable, schedule a final study visit/contact for disclosure of all remaining exam and lab test results.
* If applicable, schedule clinically indicated follow-up for subset of ongoing AEs at this visit.
* Inform the participant of planned methods and timeframes for dissemination of study results.
* Determine and document whether participant is willing to be contacted about future studies for which s/he may be eligible.
 |  |
|  | Provide reimbursement. |  |
|  | Ensure that data is entered into the study database (and perform QC2 review, if applicable) ensuring all data entered into the study database is accurate and complete.**Required Visit Forms:** * Follow-up Y/N
* Follow-up Visit Summary
* Study Discontinuation

**Log CRFs (if newly-completed or updated)*** Adverse Event Summary/Log
* Concomitant Medications Summary/Log
* Protocol Deviation Summary/Log
* Pregnancy Outcome Summary/Log (female participants only)
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**Additional Notes/Comments/Referrals:**