

OLS USE ONLY  
 [place barcode label HERE]

|                                      |  |                                      |
|--------------------------------------|--|--------------------------------------|
| <b>CT-NG, HIV, SPYHILIS, RUBELLA</b> | <b>DIAGNOSTIC IMMUNOGY LABORATORY SPECIMEN SUBMISSION FORM</b> | <b>CT-NG, HIV, SPYHILIS, RUBELLA</b> |
| <b>USE ONE FORM PER SPECIMEN</b>     |  |                                      |

| PATIENT INFORMATION   |   |                   |
|---|---|-------------------|
| PATIENT ID (Char #, MRN, etc.)  |   | Max 17 Characters |
| LAST NAME   | FIRST NAME  | MI                |
| DATE OF BIRTH   | SS# (LAST 4 DIGITS ONLY)  |                   |
| COUNTY OF RESIDENCE   | SEX (at Birth)<br><input type="checkbox"/> Female <input type="checkbox"/> Male<br><input type="checkbox"/> Not Specified                       |                   |
| STREET ADDRESS  |   |                   |
| CITY  | STATE   | ZIP               |
| PATIENT PHONE NO. (INCLUDE AREA CODE)   |   |                   |
| RACE<br><input type="checkbox"/> White <input type="checkbox"/> Asian <input type="checkbox"/> Black <input type="checkbox"/> Other<br><input type="checkbox"/> American Indian/Alaskan<br><input type="checkbox"/> Native Hawaiian or other Pacific Islander | ETHNICITY<br><input type="checkbox"/> Not Hispanic or Latino<br><input type="checkbox"/> Hispanic or Latino<br><input type="checkbox"/> Unknown |                   |
| PATIENT TYPE: <input type="checkbox"/> Employee <input type="checkbox"/> Patient <input type="checkbox"/> Investigation   |   |                   |

| DATE OF COLLECTION: |
|---------------------|
|---------------------|

| PROGRAM/CLINIC TYPE (SELECT ONE ONLY)          |   |
|--|---|
| <input type="checkbox"/> FAMILY PLANNING       | <input type="checkbox"/> STD/HIV SERVICES |
| <input type="checkbox"/> HOSPITAL              | <input type="checkbox"/> INVESTIGATION    |
| <input type="checkbox"/> CORRECTIONAL FACILITY | <input type="checkbox"/> PROJECT HRP      |
| <input type="checkbox"/> PROJECT # _____       |   |

| TEST REQUESTED (SELECT ONE ONLY)    |  |
|-------------------------------------|--|
| <input type="checkbox"/> CT/NG NAAT | <input type="checkbox"/> RUBELLA SCREEN  |
| <input type="checkbox"/> HIV PANEL  | <input type="checkbox"/> SYPHILIS SCREEN |

| SOURCE OF SPECIMEN:                   |                                      |
|---------------------------------------|--------------------------------------|
| <input type="checkbox"/> BLOOD/SERUM  | <input type="checkbox"/> THROAT SWAB |
| <input type="checkbox"/> URINE        | <input type="checkbox"/> RECTAL SWAB |
| <input type="checkbox"/> VAGINAL SWAB |                                      |

|   |
|---|
| <b>IS PATIENT PREGNANT?</b> <input type="checkbox"/> N/A <input type="checkbox"/> NO <input type="checkbox"/> YES: DUE DATE _____ |
|---|

| SUBMITTER INFORMATION   |       |     |
|---|-------|-----|
| The information below is for the mailing or faxing of test reports. Please make sure the mailing address and fax number are accurate.<br>OLS should be notified of any changes to this information as soon as possible. |       |     |
| FACILITY NAME   |       |     |
| MAILING ADDRESS   |       |     |
| CITY  | STATE | ZIP |
| COUNTY  |       |     |
| ATTENTION TO  |       |     |
| PHONE NO. (INCLUDE AREA CODE)   |       |     |
| FAX NO. (INCLUDE AREA CODE)   |       |     |

| COMPLETE FOR ALL TESTING:                     |   |
|---|---|
| <input type="checkbox"/> Routine Screen       | <input type="checkbox"/> Rescreen of previous positive (minimum of three (3) months after treatment)          |
| <input type="checkbox"/> Any symptom of STD   | <input type="checkbox"/> Suspect/possible contact to STD (new partner, multiple partners, polygamous partner) |
| <input type="checkbox"/> Known contact to STD | <input type="checkbox"/> IUD Insertion  |

| RISK FACTOR INFORMATION (SELECT ALL THAT APPLY)                         |  |
|---|--|
| PATIENT RISK FACTORS  | PARTNER RISK FACTORS   |
| <input type="checkbox"/> Sex with female                                | <input type="checkbox"/> Bisexual female   |
| <input type="checkbox"/> Sex with male                                  | <input type="checkbox"/> Bisexual male   |
| <input type="checkbox"/> Injected non-RX drugs                          | <input type="checkbox"/> IV injection drug user  |
| <input type="checkbox"/> Rec'd clotting Factor F VII A and/or F IX B    | <input type="checkbox"/> Person with AIDS or documented HIV+   |
| <input type="checkbox"/> Received transplant or artificial insemination | <input type="checkbox"/> Transfusion recipient WITH documented HIV+ and/or transplant WITH documented HIV+ |
| <input type="checkbox"/> Blood transfusion                              | <input type="checkbox"/> Person with hemophilia/clotting disorder  |
| <input type="checkbox"/> Healthcare worker and/or laboratory worker     | <b>PLACE HIV TEST FORM BARCODE LABEL HERE</b>  |
| <input type="checkbox"/> Unspecified Risk(s)                            |  |

| Provider and /or OLS Notes:   |  |
|---|--|
|   |  |
| <b>OLS USE ONLY</b><br><input type="checkbox"/> UNSAT<br>REASON/ID: | <b>ACC:</b><br><b>DE:</b><br><b>CKD:</b> |

**NOTE: RISK FACTORS FOR NO CHARGE TESTING MUST BE MARKED ON THE FORM AT THE TIME THE SPECIMEN IS RECEIVED FOR EACH REQUESTED TEST.**