# Test Order

## Lymphogranuloma venereum (LGV) Real-time PCR

<table>
<thead>
<tr>
<th>Synonym(s)</th>
<th>LGV PCR</th>
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<tr>
<td>Methodology</td>
<td>Real-time polymerase chain reaction (PCR)</td>
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<tr>
<td>Acceptable Specimen Type(s) for Testing</td>
<td>Rectal and genital swabs.</td>
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<tr>
<td>Transport / Collection Medium</td>
<td>Universal (or viral) transport medium (UTM or VTM) w/ appropriate APTIMA Collection Kit for chlamydia NAAT testing prior to LGV PCR testing. See sources below.</td>
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**Aptima® Unisex Swab Specimen Collection Kit** (catalog # 301041):
Urethral specimens only

**Aptima® Multitest Swab Specimen Collection Kit** (catalog # PRD-03546):
for rectal and vaginal specimens.

**Alinity m multi-collect specimen collection kit:** rectal and vaginal specimens.

If either Aptima® Unisex or Multitest or Alinity multi-collect Collection kit is submitted:
Please indicate whether Chlamydia NAAT screening has already been performed on specimen and date of detection. If requesting CT screening in addition to LGV PCR, please indicate both tests on requisition.

**Storage and Preservation of Specimen**

- **UTM or VTM**: Store at 2-8°C for up to 72 h; otherwise freeze at or below -30°C.
- **APTIMA collection kits**: Store at room temperature for up to 60 days.
- **Alinity m Multi-collect kits**: specimens: Store at 2-30°C for 14 days and at -25°C to -15°C for longer storage. (Avoid more than 4 freeze thaw cycles).

**Minimum Volume Required**

- 2 mL Aptima Collection kits
- 500µl Alinity m Multi-collect collection kits.

**Additional Collection Instructions**

**Additional Required Information**

Collection date required. Date of Chlamydia-positive detection

**Send Out?**

No

**Turnaround Time**

7 business days from receipt

**Testing Restrictions**

The LGV PCR test will only be performed if the NAAT CT result is positive and the LGV PCR specimen is the same specimen or was collected on the same day as the NAAT collection. If it is requested that SFDPH Laboratory perform NAAT testing for Chlamydia diagnosis or to confirm Chlamydia infection, submit requisition with appropriate tests indicated.

**Requisition Form(s)**


**Limitations / Notes / Disclaimers**

This PCR assay was developed and its performance characteristics were determined by the San Francisco Public Health Laboratory. It has not been cleared or approved by the U.S. Food and Drug Administration

Updated 11/14/23