

Reporting LGV

Serovars L₁, L₂, and L₃ of *Chlamydia trachomatis* cause a specific type of chlamydial infection, known as Lymphogranuloma Venereum (LGV). When serovars L₁, L₂, and L₃ are detected by a laboratory test, LGV is reportable as a type of *Chlamydia trachomatis*. Laboratory-confirmed cases of *Chlamydia trachomatis* infections (including serotypes L₁, L₂, and L₃) must be reported to MDH within one working day. To report LGV to the Minnesota Department of Health/MDH, please call 651-201-5414.

For further information regarding LGV diagnosis and treatment, please see [CDC: Lymphogranuloma Venereum \(LGV\) \(www.cdc.gov/std/treatment-guidelines/lgv.htm\)](https://www.cdc.gov/std/treatment-guidelines/lgv.htm).

The following reference laboratory performs LGV testing which detect serovars L₁, L₂, and L₃. Please consult the laboratory directly for additional information including specimen submission and transport requirements.

LGV Test Information

- **Laboratory:** ARUP Laboratories
- **Test code:** 2013768
- **Test Description:** *Chlamydia trachomatis* L serovars (LGV) by PCR

Minnesota Department of Health
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To obtain this information in a different format, call: 651-201-5414.