

K851033 SYNCORTECH CHLAMYDIA DIRECT SPECIMAN KITJun 21, 1985
101 days to decisionK851033 · Product code: **LJP** · Microbiology
Source: <https://www.510kdatabase.net/k851033/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antiserum, Fluorescent, Chlamydia Trachomatis (LJP)
Date received	Mar 12, 1985
Decision date	Jun 21, 1985
Days to decision	101 days
Third-party review	No

APPLICANT

Company	Syncor Intl. Corp.
Location	Mchenry, IL, US
Contact	ELEANOR V CHIU
510(k) history	31 submissions · 31 cleared · 1983-1995

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k851033/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026