FDA 510(k) Premarket Notification

K851033 SYNCORTECH CHLAMYDIA DIRECT SPECIMAN KIT

CLEARED

Jun 21, 1985
101 days to decision

K851033 · 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