FDA 510(k) Premarket Notification

K862320 DIFCO CHLAMYDIA DIRECT DETECTION SYSTEM

CLEARED

Aug 19, 1986
62 days to decision

K862320 · Product code: **LJP** · Microbiology
Source: https://www.510kdatabase.net/k862320/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antiserum, Fluorescent, Chlamydia Trachomatis (LJP)
Date received	Jun 18, 1986
Decision date	Aug 19, 1986
Days to decision	62 days
Third-party review	No

APPLICANT

Company	**Difco Laboratories, Inc.**
Location	Mchenry, IL, US
Contact	WALTER S FISHER
510(k) history	121 submissions · 121 cleared · 1979-1997

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k862320/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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