

K881500 ORTHO* CHLAMYDIA DIRECT DETECTION (FA) TESTApr 27, 1988
16 days to decisionK881500 · Product code: **LJP** · Microbiology
Source: <https://www.510kdatabase.net/k881500/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antiserum, Fluorescent, Chlamydia Trachomatis (LJP)
Date received	Apr 11, 1988
Decision date	Apr 27, 1988
Days to decision	16 days
Third-party review	No

APPLICANT

Company	Ortho Diagnostic Systems, Inc.
Location	Carpinteria, CA, US
Contact	LARRY D MCCLAIN,PHD
510(k) history	126 submissions · 126 cleared · 1981-1997

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