

**K900870 PATHODX CHLAMYDIA TRACHOMATIS DIRECT  
SPECIMEN TEST**Apr 16, 1990  
51 days to decisionK900870 · Product code: **LJP** · Microbiology  
Source: <https://www.510kdatabase.net/k900870/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antiserum, Fluorescent, Chlamydia Trachomatis (LJP)
Date received	Feb 24, 1990
Decision date	Apr 16, 1990
Days to decision	51 days
Third-party review	No
Combination product	No
PCCP authorized	No

**APPLICANT**

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Company	<b>Diagnostic Products Corp.</b>
Location	Mchenry, IL, US
Contact	KENNETH B ASARCH
510(k) history	321 submissions · 321 cleared · 1976-2006

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