

K830342 CULUTURE SET CHLAMYDIA ISOLATION ANDMay 4, 1983
92 days to decisionK830342 · Product code: **LKI** · Microbiology
Source: <https://www.510kdatabase.net/k830342/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antisera, Fluorescent, Chlamydia Spp. (LKI)
Date received	Feb 1, 1983
Decision date	May 4, 1983
Days to decision	92 days
Third-party review	No

APPLICANT

Company	ImmuloK, Inc.
Location	Mchenry, IL, US
510(k) history	15 submissions · 15 cleared · 1981-1983

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Device record: <https://www.510kdatabase.net/k830342/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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