

K811225 HUMAN BLOOD IN ALSEVER'SJun 12, 1981
39 days to decisionK811225 · Product code: **JSK** · Pathology
Source: <https://www.510kdatabase.net/k811225/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Supplement, Culture Media (JSK)
Date received	May 4, 1981
Decision date	Jun 12, 1981
Days to decision	39 days
Third-party review	No

APPLICANT

Company	Dutchland Laboratories, Inc.
Location	Mchenry, IL, US
510(k) history	75 submissions · 75 cleared · 1981-1982

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Device record: <https://www.510kdatabase.net/k811225/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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