FDA 510(k) Premarket Notification

K811225 HUMAN BLOOD IN ALSEVER'S

CLEARED

Jun 12, 1981
39 days to decision

K811225 · 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