

K791283 ANTI-HUMAN LAMBDA SERUMNov 27, 1979
140 days to decisionK791283 · Product code: **DEH** · Immunology
Source: <https://www.510kdatabase.net/k791283/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lambda, Antigen, Antiserum, Control (DEH)
Date received	Jul 10, 1979
Decision date	Nov 27, 1979
Days to decision	140 days
Third-party review	No

APPLICANT

Company	Diagnostica, Inc.
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1978-2001

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k791283/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 28, 2026