

K951101 LEUKO-TESTApr 24, 1995
45 days to decisionK951101 · Product code: **DEG** · Immunology
Source: <https://www.510kdatabase.net/k951101/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lactoferrin, Antigen, Antiserum, Control (DEG)
Date received	Mar 10, 1995
Decision date	Apr 24, 1995
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Techlab, Inc.
Location	Blacksburg, VA, US
Contact	DAVID M LYERLY
Website	http://www.techlab.com/
510(k) history	36 submissions · 36 cleared · 1992-2019

Techlab, Inc. designs, develops, and manufactures infectious disease diagnostics in the USA. The company specializes in enteric and microbiology diagnostic products distributed worldwide. Techlab holds ISO 13485 certification and MDSAP approval with FDA registration. Techlab received FDA 510(k) clearances from total submissions between 1992 and 2019. The company's portfolio is dominated by microbiology devices, representing 83% of submissions. Notable cleared products include tests for *Clostridioides difficile*, *Helicobacter pylori*, parasites, and fecal biomarkers. Techlab...
