

K831365 TESTCORP ANAJun 2, 1983
37 days to decisionK831365 · Product code: **LKS** · Microbiology
Source: <https://www.510kdatabase.net/k831365/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Parasite Concentration (LKS)
Date received	Apr 26, 1983
Decision date	Jun 2, 1983
Days to decision	37 days
Third-party review	No

APPLICANT

Company	Catheter Technology Corp.
Location	Walker, MI, US
510(k) history	13 submissions · 13 cleared · 1983-1988

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k831365/> 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 July 1, 2026