FDA 510(k) Premarket Notification

K831365 TESTCORP ANA

CLEARED

Jun 2, 1983
37 days to decision

K831365 · 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