

K954647 INCLINOMETEROct 30, 1995
20 days to decisionK954647 · Product code: **KQX** · Neurology
Source: <https://www.510kdatabase.net/k954647/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Goniometer, Ac-powered (KQX)
Date received	Oct 10, 1995
Decision date	Oct 30, 1995
Days to decision	20 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	J-Tech, Inc.
Location	Midvale, UT, US
Contact	J. TRACY LIVINGSTON
510(k) history	5 submissions · 5 cleared · 1991-1996

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k954647/> 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 30, 2026