

K962377 3D-SPINESep 3, 1996
75 days to decisionK962377 · Product code: **KQX** · Neurology
Source: <https://www.510kdatabase.net/k962377/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Goniometer, Ac-powered (KQX)
Date received	Jun 20, 1996
Decision date	Sep 3, 1996
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Skill Technologies, Inc.
Location	Phoenix, AZ, US
510(k) history	1 submissions · 1 cleared · 1996-1996

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k962377/>, 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