FDA 510(k) Premarket Notification

K962377 3D-SPINE

CLEARED

Sep 3, 1996
75 days to decision

K962377 · 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