FDA 510(k) Premarket Notification

K894283 COMPREHENSIVE SPINE MOTION ANALYZER

CLEARED

Jan 19, 1990
210 days to decision

K894283 · Product code: **KQX** · Neurology
Source: https://www.510kdatabase.net/k894283/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Goniometer, Ac-powered (KQX)
Date received	Jun 23, 1989
Decision date	Jan 19, 1990
Days to decision	210 days
Third-party review	No

APPLICANT

Company	**Orthopedic Systems, Inc.**
Location	Mchenry, IL, US
Contact	ROBERT MOORE
510(k) history	95 submissions · 89 cleared · 1977-1996