FDA 510(k) Premarket Notification

K914060 RANGE OF MOTION SENSOR, MODEL RM001

CLEARED

Dec 9, 1991
90 days to decision

K914060 · Product code: **KQX** · Neurology
Source: https://www.510kdatabase.net/k914060/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Goniometer, Ac-powered (KQX)
Date received	Sep 10, 1991
Decision date	Dec 9, 1991
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Nk Biotechnical Engineering Co.**
Location	Minneapolis, MN, US
Contact	KAREN GOTFREDSON
510(k) history	15 submissions · 15 cleared · 1991-1994