

K914060 RANGE OF MOTION SENSOR, MODEL RM001Dec 9, 1991
90 days to decisionK914060 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k914060/> 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 May 22, 2026