

**K943929 DATAGLOVE AND MOVEMENT ANALYSIS SYSTEM
(MAS)**May 15, 1995
276 days to decisionK943929 · Product code: **KQX** · Neurology
Source: <https://www.510kdatabase.net/k943929/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Goniometer, Ac-powered (KQX)
Date received	Aug 12, 1994
Decision date	May 15, 1995
Days to decision	276 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Greenleaf Medical Systems, Inc.
Location	Palo Alto, CA, US
Contact	GLENN R EDWARDS
510(k) history	4 submissions · 4 cleared · 1989-1995

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k943929/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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