

**K884723 (DATAGLOVE) HAND IMPAIRMENT EVALUATION
SYSTEM**Feb 8, 1989
90 days to decisionK884723 · Product code: **KQX** · Neurology
Source: <https://www.510kdatabase.net/k884723/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Goniometer, Ac-powered (KQX)
Date received	Nov 10, 1988
Decision date	Feb 8, 1989
Days to decision	90 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k884723/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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