FDA 510(k) Premarket Notification

K884723 (DATAGLOVE) HAND IMPAIRMENT EVALUATION SYSTEM

CLEARED

Feb 8, 1989
90 days to decision

K884723 · 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