FDA 510(k) Premarket Notification

K943929 DATAGLOVE AND MOVEMENT ANALYSIS SYSTEM (MAS)

CLEARED

May 15, 1995
276 days to decision

K943929 · 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