FDA 510(k) Premarket Notification

K893537 GREENLEAF HAND IMPAIRMENT EVALUATION SYSTEM

CLEARED

Oct 10, 1989
155 days to decision

K893537 · Product code: **LBB** · Neurology
Source: https://www.510kdatabase.net/k893537/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dynamometer, Ac-powered (LBB)
Date received	May 8, 1989
Decision date	Oct 10, 1989
Days to decision	155 days
Third-party review	No

APPLICANT

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