

**K893537 GREENLEAF HAND IMPAIRMENT EVALUATION SYSTEM**Oct 10, 1989  
155 days to decisionK893537 · Product code: **LBB** · Neurology  
Source: <https://www.510kdatabase.net/k893537/>**SUBMISSION DETAILS**

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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**

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

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k893537/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 1, 2026