

**K831550 FINGER BONIOMETER BK7506**Jun 24, 1983  
43 days to decisionK831550 · Product code: **KQW** · Neurology  
Source: <https://www.510kdatabase.net/k831550/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Goniometer, Nonpowered (KQW)
Date received	May 12, 1983
Decision date	Jun 24, 1983
Days to decision	43 days
Third-party review	No

**APPLICANT**

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Company	<b>Fred Sammons, Inc.</b>
Location	Mchenry, IL, US
Website	<a href="https://www.sammons-preston.com">https://www.sammons-preston.com</a>
510(k) history	278 submissions · 278 cleared · 1976-1988

Fred Sammons, Inc. is a medical device company based in McHenry, US, specializing in rehabilitation products and assistive devices for patient care and therapy. The company has received FDA 510(k) clearances from total submissions, with 88% focused on Physical Medicine devices. FDA 510(k) clearances span from 1976 to 1988, establishing a historical regulatory record in therapeutic aids, orthotic supports, and mobility assistance equipment. Notable cleared devices include orthotic plastics, arm slings, splints, traction exercise equipment, and specialized assessment tools....

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k831550/> 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 May 28, 2026