

**K801717 BK-7511 BRASS GONIOMETER**Aug 4, 1980  
11 days to decisionK801717 · Product code: **KQW** · Neurology  
Source: <https://www.510kdatabase.net/k801717/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Goniometer, Nonpowered (KQW)
Date received	Jul 24, 1980
Decision date	Aug 4, 1980
Days to decision	11 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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