FDA 510(k) Premarket Notification

# K810441 UNIVERSAL GONIMETER

**CLEARED**

Feb 27, 1981
8 days to decision

K810441 · Product code: **KQW** · Neurology
Source: https://www.510kdatabase.net/k810441/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Goniometer, Nonpowered (KQW) |
| Date received | Feb 19, 1981 |
| Decision date | Feb 27, 1981 |
| Days to decision | 8 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Fred Sammons, Inc.** |
| Location | Mchenry, IL, US |
| Website | https://www.sammons-preston.com |
| 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....

**510k Database** - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k810441/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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