FDA 510(k) Premarket Notification

K810222 LIMB ORHTOSIS

CLEARED

Feb 4, 1981
8 days to decision

K810222 · Product code: **ITQ** · Physical Medicine
Source: https://www.510kdatabase.net/k810222/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Joint, Knee, External Brace (ITQ)
Date received	Jan 27, 1981
Decision date	Feb 4, 1981
Days to decision	8 days
Third-party review	No

APPLICANT

Company	**Medical Design & Manufacturing Corp.**
Location	Walker, MI, US
510(k) history	10 submissions · 10 cleared · 1981-1992