

K810222 LIMB ORHTOSISFeb 4, 1981
8 days to decisionK810222 · Product code: **ITQ** · Physical MedicineSource: <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

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Device record: <https://www.510kdatabase.net/k810222/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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