

K833084 MASTERHINGEOct 14, 1983
31 days to decisionK833084 · Product code: **LGF** · Orthopedic
Source: <https://www.510kdatabase.net/k833084/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Component, Cast (LGF)
Date received	Sep 13, 1983
Decision date	Oct 14, 1983
Days to decision	31 days
Third-party review	No

APPLICANT

Company	Mark One Healthcare Products
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1977-1989

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Device record: <https://www.510kdatabase.net/k833084/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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