FDA 510(k) Premarket Notification

K833084 MASTERHINGE

CLEARED

Oct 14, 1983
31 days to decision

K833084 · 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