FDA 510(k) Premarket Notification

K834384 MODULAR EXTREMITY CASTING SUPPORT

CLEARED

Jan 30, 1984
47 days to decision

K834384 · Product code: **LGG** · Orthopedic
Source: https://www.510kdatabase.net/k834384/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Cast Application/removal, Manual (LGG)
Date received	Dec 14, 1983
Decision date	Jan 30, 1984
Days to decision	47 days
Third-party review	No

APPLICANT

Company	**Buckman Co., Inc.**
Location	Mchenry, IL, US
510(k) history	111 submissions · 104 cleared · 1983-1998