

K834384 MODULAR EXTREMITY CASTING SUPPORTJan 30, 1984
47 days to decisionK834384 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k834384/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026