

K003128 MODIFICATION TO RELIEF BRIEFAug 12, 2002
675 days to decisionK003128 · Product code: **NJB** · Physical Medicine
Source: <https://www.510kdatabase.net/k003128/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Truncal, For Dysmenorrhea (NJB)
Date received	Oct 6, 2000
Decision date	Aug 12, 2002
Days to decision	675 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	The JM Kohn Co.
Location	San Francisco, CA, US
Contact	CHARLES L MORIN
510(k) history	1 submissions · 1 cleared · 2002-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k003128/> 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 June 29, 2026