FDA 510(k) Premarket Notification

K003128 MODIFICATION TO RELIEF BRIEF

CLEARED

Aug 12, 2002
675 days to decision

K003128 · 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