FDA 510(k) Premarket Notification

K812342 ZEE EXTRICATION DEVICE

CLEARED

Sep 21, 1981
34 days to decision

K812342 · Product code: **IQF** · Physical Medicine
Source: https://www.510kdatabase.net/k812342/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Cervical-thoracic, Rigid (IQF)
Date received	Aug 18, 1981
Decision date	Sep 21, 1981
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Zee Medical Products Co., Inc.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1981-1983

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k812342/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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