FDA 510(k) Premarket Notification

K823105 RESI-WALKER

CLEARED

Nov 5, 1982
16 days to decision

K823105 · Product code: **ITJ** · Physical Medicine
Source: https://www.510kdatabase.net/k823105/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Walker, Mechanical (ITJ)
Date received	Oct 20, 1982
Decision date	Nov 5, 1982
Days to decision	16 days
Third-party review	No

APPLICANT

Company	**Orthion Corp.**
Location	Mchenry, IL, US
510(k) history	11 submissions · 11 cleared · 1981-1983