FDA 510(k) Premarket Notification

K905113 ROTO REST

CLEARED

Nov 26, 1990
13 days to decision

K905113 · Product code: **IKZ** · Physical Medicine
Source: https://www.510kdatabase.net/k905113/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bed, Patient Rotation, Powered (IKZ)
Date received	Nov 13, 1990
Decision date	Nov 26, 1990
Days to decision	13 days
Third-party review	No
Other names	TRAUMA AND INTENSIVE CARE BED

APPLICANT

Company	**Ethos Medical Products , Ltd.**
Location	Athlone, IE
Contact	PAT CONNOLLY
510(k) history	1 submissions · 1 cleared · 1990-1990