FDA 510(k) Premarket Notification

K905756 TITAN MERIDIAN INTERSEGMENTAL ROLLERTABLE

CLEARED

Jan 8, 1991
12 days to decision

K905756 · Product code: **JFB** · Physical Medicine
Source: https://www.510kdatabase.net/k905756/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Physical Therapy, Multi Function (JFB)
Date received	Dec 27, 1990
Decision date	Jan 8, 1991
Days to decision	12 days
Third-party review	No

APPLICANT

Company	**Titan Technology Intl., Inc.**
Location	St. Louis, MI, US
Contact	HOWARD COOPER
510(k) history	1 submissions · 1 cleared · 1991-1991