FDA 510(k) Premarket Notification

K832940 ROTAGYM

CLEARED

Nov 7, 1983
70 days to decision

K832940 · Product code: **ISD** · Physical Medicine
Source: https://www.510kdatabase.net/k832940/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Exerciser, Measuring (ISD)
Date received	Aug 29, 1983
Decision date	Nov 7, 1983
Days to decision	70 days
Third-party review	No

APPLICANT

Company	**I-Rep, Inc.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1982-1994