

K832940 ROTAGYMNov 7, 1983
70 days to decisionK832940 · Product code: **ISD** · Physical MedicineSource: <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

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Device record: <https://www.510kdatabase.net/k832940/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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