FDA 510(k) Premarket Notification

K812902 ACUPRESS VIBRATOR

CLEARED

Nov 10, 1981
25 days to decision

K812902 · Product code: **IRO** · Physical Medicine
Source: https://www.510kdatabase.net/k812902/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Vibrator, Therapeutic (IRO)
Date received	Oct 16, 1981
Decision date	Nov 10, 1981
Days to decision	25 days
Third-party review	No

APPLICANT

Company	**Eicona Corp.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981