FDA 510(k) Premarket Notification

K823381 PROFESSIONAL MARK I BIOFEEDBACK SYS

CLEARED

Jan 5, 1983
54 days to decision

K823381 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k823381/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Nov 12, 1982
Decision date	Jan 5, 1983
Days to decision	54 days
Third-party review	No

APPLICANT

Company	**Srs Co.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1982-1983