

K823381 PROFESSIONAL MARK I BIOFEEDBACK SYSJan 5, 1983
54 days to decisionK823381 · 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

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Device record: <https://www.510kdatabase.net/k823381/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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