FDA 510(k) Premarket Notification

K812179 LIFE-TECH #1753 BIOFEEDBACK TRAINER

CLEARED

Aug 25, 1981
22 days to decision

K812179 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k812179/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Aug 3, 1981
Decision date	Aug 25, 1981
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Life-Tech Instruments, Inc.**
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1977-1981

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k812179/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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