FDA 510(k) Premarket Notification

K834110 ANTENSE ANTI-TENSION DEVICE

CLEARED

Feb 4, 1984
67 days to decision

K834110 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k834110/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Nov 29, 1983
Decision date	Feb 4, 1984
Days to decision	67 days
Third-party review	No

APPLICANT

Company	**Bio Sig Instruments, Inc.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1978-1984