

**K834110 ANTENSE ANTI-TENSION DEVICE**Feb 4, 1984  
67 days to decisionK834110 · Product code: **HCC** · Neurology  
Source: <https://www.510kdatabase.net/k834110/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k834110/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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