

**K884852 BIOFEEDBACK DEVICE EMG1, EMG ULTRA LOW NOISE OPTI.**Feb 2, 1989  
76 days to decisionK884852 · Product code: **HCC** · Neurology  
Source: <https://www.510kdatabase.net/k884852/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Nov 18, 1988
Decision date	Feb 2, 1989
Days to decision	76 days
Third-party review	No

**APPLICANT**

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Company	<b>Moe</b>
Location	Soquel, CA, US
Contact	MIKE WILBER
510(k) history	6 submissions · 6 cleared · 1989-1989

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k884852/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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