

**K960885 MODEL 1240, SMALL E NET**Apr 26, 1996  
53 days to decisionK960885 · Product code: **GXY** · Neurology  
Source: <https://www.510kdatabase.net/k960885/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Mar 4, 1996
Decision date	Apr 26, 1996
Days to decision	53 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Physiometrix, Inc.</b>
Location	Sunnyvale, CA, US
Contact	DAWN E FRAZER
510(k) history	22 submissions · 22 cleared · 1992-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960885/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 30, 2026