

**K030800 ELECTROMED REUSABLE NEUROSTIMULATION  
ELECTRODES**Jun 11, 2003  
90 days to decisionK030800 · Product code: **GXY** · Neurology  
Source: <https://www.510kdatabase.net/k030800/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Mar 13, 2003
Decision date	Jun 11, 2003
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Eletromed, Inc.</b>
Location	Betheda, MD, US
Contact	LYDIA BAYNES
510(k) history	1 submissions · 1 cleared · 2003-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030800/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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