

**K010638 NEUROPLUS, MODEL A10040, A10041, A10042,  
A10043**May 3, 2001  
59 days to decisionK010638 · Product code: **GXY** · Neurology  
Source: <https://www.510kdatabase.net/k010638/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Mar 5, 2001
Decision date	May 3, 2001
Days to decision	59 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Vermont Medical, Inc.</b>
Location	Bellows Falls, VT, US
Contact	MARC FILLION
510(k) history	9 submissions · 9 cleared · 1978-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k010638/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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