

**K903554 NORTECH MODEL 87202204 ELECTRODE**Aug 30, 1990  
23 days to decisionK903554 · Product code: **GXY** · Neurology  
Source: <https://www.510kdatabase.net/k903554/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Aug 7, 1990
Decision date	Aug 30, 1990
Days to decision	23 days
Third-party review	No

**APPLICANT**

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Company	<b>Medtronic Andover Medical, Inc.</b>
Location	Lowell, MA, US
Contact	JANICE M PEVIDE
510(k) history	6 submissions · 6 cleared · 1985-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903554/>; 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 May 14, 2026