

**K790316 ELECTRODE, ELECTROFLEX**Mar 27, 1979  
40 days to decisionK790316 · Product code: **GXY** · Neurology  
Source: <https://www.510kdatabase.net/k790316/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Feb 15, 1979
Decision date	Mar 27, 1979
Days to decision	40 days
Third-party review	No

**APPLICANT**

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Company	<b>Andover Medical, Inc.</b>
Location	Mchenry, IL, US
Website	<a href="https://www.andovermedical.com">https://www.andovermedical.com</a>
510(k) history	21 submissions · 21 cleared · 1976-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k790316/>; 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 27, 2026