

**K862028 ULTRAHESIVE II POSTOP ELECTRODE**Aug 1, 1986  
65 days to decisionK862028 · Product code: **GXY** · Neurology  
Source: <https://www.510kdatabase.net/k862028/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	May 28, 1986
Decision date	Aug 1, 1986
Days to decision	65 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Devices, Inc.</b>
Location	Mchenry, IL, US
Contact	BRUCE MACFARLANE,PHD
510(k) history	49 submissions · 47 cleared · 1977-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k862028/>; 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 21, 2026