

**K903402 GEOHESIVE (TM), MODEL NUMBERS 1022L, 1023L,  
1026L**Aug 22, 1990  
22 days to decisionK903402 · Product code: **GXY** · Neurology  
Source: <https://www.510kdatabase.net/k903402/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Jul 31, 1990
Decision date	Aug 22, 1990
Days to decision	22 days
Third-party review	No

**APPLICANT**

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Company	<b>Medi-Source, Inc.</b>
Location	Syosset, NY, US
Contact	BRUCE MACFARLANE
510(k) history	4 submissions · 4 cleared · 1989-1990

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903402/> 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