

**K780998 SCALPEL**Jul 27, 1978  
43 days to decisionK780998 · Product code: **GDX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k780998/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Scalpel, One-piece (GDX)
Date received	Jun 14, 1978
Decision date	Jul 27, 1978
Days to decision	43 days
Third-party review	No

**APPLICANT**

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Company	<b>Isolab, Inc.</b>
Location	Mchenry, IL, US
510(k) history	44 submissions · 44 cleared · 1977-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k780998/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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