

**K880377 NOVO VIVO MODEL #17140 AND #17135**Apr 4, 1988  
68 days to decisionK880377 · Product code: **GAA** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k880377/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Aspiration And Injection, Disposable (GAA)
Date received	Jan 27, 1988
Decision date	Apr 4, 1988
Days to decision	68 days
Third-party review	No

**APPLICANT**

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Company	<b>Genesis Industries, Inc.</b>
Location	Walker, MI, US
Contact	ANDERSON, DVM
510(k) history	17 submissions · 17 cleared · 1983-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880377/>, 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 June 29, 2026