

**K890722 ADG BONE BIOPSY SYSTEM**Sep 21, 1989  
220 days to decisionK890722 · Product code: **GDM** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k890722/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Aspiration And Injection, Reusable (GDM)
Date received	Feb 13, 1989
Decision date	Sep 21, 1989
Days to decision	220 days
Third-party review	No

**APPLICANT**

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Company	<b>American Design Group, Inc.</b>
Location	Martinez, CA, US
Contact	DAVID W SCHLERF
510(k) history	3 submissions · 3 cleared · 1989-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890722/> 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 22, 2026