

**K923872 BONE MARROW BIOPSY SYSTEM**Dec 15, 1992  
134 days to decisionK923872 · Product code: **GDM** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k923872/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Aspiration And Injection, Reusable (GDM)
Date received	Aug 3, 1992
Decision date	Dec 15, 1992
Days to decision	134 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>The Straumann Co.</b>
Location	Cambridge, MA, US
Contact	CAROLYN BITETTI
510(k) history	14 submissions · 14 cleared · 1992-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k923872/> 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 July 1, 2026