

**K123297 COMPRESS SEGMENTAL HUMERAL REPLACEMENT SYSTEM**Nov 20, 2012  
29 days to decisionK123297 · Product code: **MBF** · Orthopedic  
Source: <https://www.510kdatabase.net/k123297/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Prosthesis, Shoulder, Semi-constrained, Metal/polymer, Uncemented (MBF)
Date received	Oct 22, 2012
Decision date	Nov 20, 2012
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Biomet Manufacturing Corp.</b>
Location	Warsaw, IN, US
Contact	PATRICIA BERES
510(k) history	2 submissions · 2 cleared · 2012-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k123297/>. 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 17, 2026