

**K172502 Comprehensive Augmented Glenoid Components,  
Comprehensive Standard Baseplate, Comprehensive Mini  
Baseplate**Jan 3, 2018  
138 days to decisionK172502 · Product code: **PHX** · Orthopedic  
Source: <https://www.510kdatabase.net/k172502/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shoulder Prosthesis, Reverse Configuration (PHX)
Date received	Aug 18, 2017
Decision date	Jan 3, 2018
Days to decision	138 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 Sandborn Beres
510(k) history	93 submissions · 93 cleared · 2004-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k172502/> 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 14, 2026