

**K131353 COMPREHENSIVE REVERSE SHOULDER - TITANIUM
GLENOSPHERE**Oct 8, 2013
151 days to decisionK131353 · Product code: **PHX** · Orthopedic
Source: <https://www.510kdatabase.net/k131353/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shoulder Prosthesis, Reverse Configuration (PHX)
Date received	May 10, 2013
Decision date	Oct 8, 2013
Days to decision	151 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Biomet Manufacturing Corp
Location	Warsaw, IN, US
Contact	PATRICIA S BERES
510(k) history	93 submissions · 93 cleared · 2004-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k131353/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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