

**K121183 COMPREHENSIVE REVERSE SHOULDER - E1
POLYETHYLENE CLAIMS**Jul 26, 2012
99 days to decisionK121183 · Product code: **PHX** · Orthopedic
Source: <https://www.510kdatabase.net/k121183/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shoulder Prosthesis, Reverse Configuration (PHX)
Date received	Apr 18, 2012
Decision date	Jul 26, 2012
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Biomet Corporation
Location	Warsaw, IN, US
Contact	PATRICIA SANDBORN BERES
510(k) history	1 submissions · 1 cleared · 2012-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k121183/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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