K172502 Comprehensive Augmented Glenoid Components, Comprehensive Standard Baseplate, Comprehensive Mini Baseplate

CLEARED

Jan 3, 2018
138 days to decision

K172502 · Product code: **PHX** · Orthopedic
Source: https://www.510kdatabase.net/k172502/

SUBMISSION DETAILS

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

Company	**Biomet Manufacturing Corp**
Location	Warsaw, IN, US
Contact	Patricia Sandborn Beres
510(k) history	93 submissions · 93 cleared · 2004-2023