FDA 510(k) Premarket Notification

K052888 POROUS TITANIUM ACETABULAR AUGMENTS

CLEARED

Dec 6, 2005
54 days to decision

K052888 · Product code: **KWA** · Orthopedic
Source: https://www.510kdatabase.net/k052888/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained (metal Uncemented Acetabular Component) (KWA)
Date received	Oct 13, 2005
Decision date	Dec 6, 2005
Days to decision	54 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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