

K052888 POROUS TITANIUM ACETABULAR AUGMENTSDec 6, 2005
54 days to decisionK052888 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k052888/> Data sourced from FDA 510(k) public records (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. - Generated May 20, 2026