

**K921139 HEALEY REVISION COMPONENT SYSTEM**Jul 1, 1992  
113 days to decisionK921139 · Product code: **JDL** · Orthopedic  
Source: <https://www.510kdatabase.net/k921139/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained (metal Cemented Acetabular Component) (JDL)
Date received	Mar 10, 1992
Decision date	Jul 1, 1992
Days to decision	113 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Biomet, Inc.</b>
Location	McHenry, IL, US
Contact	DUTCH HARRISON
Website	<a href="http://www.biomet.com/">http://www.biomet.com/</a>
510(k) history	440 submissions · 418 cleared · 1978-2024

Biomet, Inc. is an orthopedic medical device manufacturer based in McHenry, US. The company specializes in surgical implants, fixation systems, and trauma solutions. Biomet has maintained a strong FDA 510(k) regulatory record since its first clearance in 1978. The company has received FDA 510(k) clearances from total submissions. Orthopedic devices represent 88% of its submission portfolio, reflecting the company's core focus on joint reconstruction, trauma fixation, and surgical instrumentation. The latest clearance in 2024 demonstrates continued regulatory activity and ...