

K001374 PATIENT MATCHED COBALT ALLOY TROCHLEA REPLACEMENTJul 27, 2000
87 days to decisionK001374 · Product code: **KRR** · Orthopedic
Source: <https://www.510kdatabase.net/k001374/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patello/femoral, Semi-constrained, Cemented, Metal/polymer (KRR)
Date received	May 1, 2000
Decision date	Jul 27, 2000
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Biomet Manufacturing, Inc.
Location	Warsaw, IN, US
Contact	PATRICIA SANDBORN BERES
510(k) history	32 submissions · 32 cleared · 1999-2013

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