

**K002356 NATURAL-KNEE II PATELLOFEMORAL JOINT  
PROSTHESIS**Oct 30, 2000  
89 days to decisionK002356 · Product code: **KRR** · Orthopedic  
Source: <https://www.510kdatabase.net/k002356/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patello/femoral, Semi-constrained, Cemented, Metal/polymer (KRR)
Date received	Aug 2, 2000
Decision date	Oct 30, 2000
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sulzer Orthopedics, Inc.</b>
Location	Austin, TX, US
Contact	MITCHELL A DHORITY
510(k) history	45 submissions · 41 cleared · 1997-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k002356/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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