

**K023667 RT-PLUS SOLUTION & RT-PLUS MODULAR SOLUTION KNEE**

Dec 24, 2002  
54 days to decision

K023667 · Product code: **KRO** · Orthopedic  
Source: <https://www.510kdatabase.net/k023667/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Prosthesis, Knee, Femorotibial, Constrained, Cemented, Metal/polymer (KRO)
Date received	Oct 31, 2002
Decision date	Dec 24, 2002
Days to decision	54 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Plus Orthopedics</b>
Location	San Diego, CA, US
Contact	J.D. WEBB
510(k) history	38 submissions · 38 cleared · 1997-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k023667/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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