FDA 510(k) Premarket Notification

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

CLEARED

Dec 24, 2002
54 days to decision

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

SUBMISSION DETAILS

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

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