

**K013982 KINEMATCH PATELLO-FEMORAL RESURFACING
IMPALANT**May 21, 2002
169 days to decisionK013982 · Product code: **KRR** · Orthopedic
Source: <https://www.510kdatabase.net/k013982/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patello/femoral, Semi-constrained, Cemented, Metal/polymer (KRR)
Date received	Dec 3, 2001
Decision date	May 21, 2002
Days to decision	169 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Kinamed, Inc.
Location	Los Angeles, CA, US
Contact	VINEET SARIN
510(k) history	33 submissions · 29 cleared · 1988-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k013982/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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