

**K092441 MODIFICATION TO: CONFORMIS IUNI UNICONDYLAR
KNEE REPAIR SYSTEM**Sep 9, 2009
30 days to decisionK092441 · Product code: **HSX** · Orthopedic
Source: <https://www.510kdatabase.net/k092441/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Prosthesis, Knee, Femorotibial, Non-constrained, Cemented, Metal/polymer (HSX)
Date received	Aug 10, 2009
Decision date	Sep 9, 2009
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Conformis, Inc.
Location	Foster City, CA, US
Contact	AMITA SHAH
510(k) history	60 submissions · 60 cleared · 2005-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k092441/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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