

**K231233 Identity™ Imprint™ CR KRS and Identity™ Imprint™ PS KRS**

May 26, 2023  
28 days to decision

K231233 · Product code: **JWH** · Orthopedic  
Source: <https://www.510kdatabase.net/k231233/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Apr 28, 2023
Decision date	May 26, 2023
Days to decision	28 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Conformis, Inc.</b>
Location	Foster City, CA, US
Contact	Nivedita Namjoshi
510(k) history	60 submissions · 60 cleared · 2005-2023

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

Device record: <https://www.510kdatabase.net/k231233/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).

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