

**K230120 KeYi Total Knee System**Apr 19, 2024  
458 days to decisionK230120 · Product code: **JWH** · Orthopedic  
Source: <https://www.510kdatabase.net/k230120/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Jan 17, 2023
Decision date	Apr 19, 2024
Days to decision	458 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Beijing Keyi Medical Device Technology Co., Ltd.</b>
Location	Beijing, CN
Contact	Jenny Jiang
510(k) history	4 submissions · 4 cleared · 2017-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k230120/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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