

K172189 Ins Hilden Humeral Arzzt, Ins Hilden Cannulated Tibial Arzzt, Ins Hilden Femoral Arzzt & Ins Hilden Retrograde Femoral Arzzt

Jan 16, 2018
180 days to decision

K172189 · Product code: **HSB** · Orthopedic
Source: <https://www.510kdatabase.net/k172189/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Rod, Fixation, Intramedullary And Accessories (HSB)
Date received	Jul 20, 2017
Decision date	Jan 16, 2018
Days to decision	180 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Truemed Group, LLC
Location	Hampton, VA, US
Contact	Nina Galeana Rodriguez
510(k) history	5 submissions · 5 cleared · 2014-2021

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k172189/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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