

K160700 ATTUNE Revision Knee System

Jun 10, 2016
88 days to decision

K160700 · Product code: **JWH** · Orthopedic
Source: <https://www.510kdatabase.net/k160700/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Mar 14, 2016
Decision date	Jun 10, 2016
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Depuy(Ireland)
Location	Cork, IE
Contact	Jaime Weeks
510(k) history	13 submissions · 13 cleared · 2010-2022

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k160700/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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