

K242665 DePuy ATTUNE™ Total Knee SystemNov 1, 2024
58 days to decisionK242665 · Product code: **OIY** · Orthopedic
Source: <https://www.510kdatabase.net/k242665/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer + Additive/metal/polymer + Additive (OIY)
Date received	Sep 4, 2024
Decision date	Nov 1, 2024
Days to decision	58 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Depuy Orthopedics, Inc.
Location	Warsaw, IN, US
Contact	Christina Kaucic
510(k) history	1 submissions · 1 cleared · 2024-2024

REGULATORY CONSULTANT

Consulting firm	Depuy Ireland UC
Contact	Susan Mullane

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k242665/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026