

K252887 DePuy ATTUNE™ Knee SystemMay 22, 2026
254 days to decisionK252887 · Product code: **OIY** · Orthopedic
Source: <https://www.510kdatabase.net/k252887/>**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 10, 2025
Decision date	May 22, 2026
Days to decision	254 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	DePuy Orthopaedics, Inc.
Location	Warsaw, IN, US
Contact	Christina Kaucic
510(k) history	207 submissions · 205 cleared · 1998-2026

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

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