

**K212933 INHANCETM Hybrid Anatomic Glenoid Implant**Jun 8, 2022  
266 days to decisionK212933 · Product code: **MBF** · Orthopedic  
Source: <https://www.510kdatabase.net/k212933/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Shoulder, Semi-constrained, Metal/polymer, Uncemented (MBF)
Date received	Sep 15, 2021
Decision date	Jun 8, 2022
Days to decision	266 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Depuy Ireland UC</b>
Location	Ringaskiddy, IE
Contact	Yayoi Fujimaki
510(k) history	47 submissions · 47 cleared · 2018-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Ignite Orthopedics, LLC</b>
Contact	Russ Parrott

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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