

K231503 CUPTIMIZE™ AdvancedAug 22, 2023
90 days to decisionK231503 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k231503/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	May 24, 2023
Decision date	Aug 22, 2023
Days to decision	90 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	DePuy Orthopaedics, Inc.
Location	Warsaw, IN, US
Contact	Sierra Robinson
510(k) history	206 submissions · 204 cleared · 1998-2023

REGULATORY CONSULTANT

Consulting firm	Depuy Ireland UC
Contact	Sierra Robinson

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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