

K220449 DePuy CONDUIT LLIF SQUID InserterJun 28, 2022
132 days to decisionK220449 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k220449/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Feb 16, 2022
Decision date	Jun 28, 2022
Days to decision	132 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Enztec Limited
Location	Middleton, NZ
Contact	Claire Mai
510(k) history	5 submissions · 5 cleared · 2021-2023

REGULATORY CONSULTANT

Consulting firm	Empirical Testing Corp
Contact	Nathan Wright

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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