

K223671 CONDUIT LLIF Straight InsertersJan 5, 2023
29 days to decisionK223671 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k223671/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 7, 2022
Decision date	Jan 5, 2023
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k223671/> 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 26, 2026