

**K172480 icotec Interbody Cage System**May 17, 2018  
274 days to decisionK172480 · Product code: **ODP** · Orthopedic  
Source: <https://www.510kdatabase.net/k172480/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Aug 16, 2017
Decision date	May 17, 2018
Days to decision	274 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Icotec AG</b>
Location	Altstaetten, SE
Contact	Valeria Consiero
510(k) history	16 submissions · 16 cleared · 2016-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k172480/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 25, 2026