

K222973 IdentiTi and Transcend Interbody Systems: IdentiTi Cervical Porous Ti Interbody System, IdentiTi NanoTec Cervical Interbody System, Transcend Cervical PEEK Interbody System, Transcend NanoTec Cervical Interbody System, IdentiTi Cervical Standalone Interbody System, IdentiTi NanoTec Cervical Standalone Interbody System

Nov 17, 2022
50 days to decision

K222973 · Product code: **ODP** · Orthopedic
Source: <https://www.510kdatabase.net/k222973/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Sep 28, 2022
Decision date	Nov 17, 2022
Days to decision	50 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Alphatec Spine, Inc.
Location	Carlsbad, CA, US
Contact	Andrew Zhang
Website	https://www.alphatecspine.com
510(k) history	93 submissions · 93 cleared · 2005-2026

Alphatec Spine, Inc. is a spine surgery medical device company based in Carlsbad, California. The company develops and markets surgical solutions for spinal fusion and fixation procedures. Alphatec Spine maintains a strong FDA 510(k) regulatory record with FDA 510(k) clearances from total submissions. The company specializes in Orthopedic devices, which represent 91% of its submission portfolio. Clearances span from 2005 to 2026, demonstrating sustained regulatory activity and recent market engagement. Recent cleared devices include robotic navigation systems, interbody s...