

K251725 ANAX™ OCT Spinal SystemJun 25, 2025
20 days to decisionK251725 · Product code: **NKG** · Orthopedic
Source: <https://www.510kdatabase.net/k251725/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Posterior Cervical Screw System (NKG)
Date received	Jun 5, 2025
Decision date	Jun 25, 2025
Days to decision	20 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Cg Medtech Co., Ltd.
Location	Uijeongbu-Si, KR
Contact	Jeongah Kim
510(k) history	2 submissions · 2 cleared · 2025-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k251725/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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