

K213871 Ennovate Cervical Spinal and Occiput SystemJul 1, 2022
200 days to decisionK213871 · Product code: **NKG** · Orthopedic
Source: <https://www.510kdatabase.net/k213871/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Posterior Cervical Screw System (NKG)
Date received	Dec 13, 2021
Decision date	Jul 1, 2022
Days to decision	200 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Aesculap Implant Systems, LLC
Location	Center Valley, PA, US
Contact	Paul Amudala
510(k) history	22 submissions · 22 cleared · 2010-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k213871/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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