

**K231769 activL® Next Generation Instrumentation**Jul 13, 2023  
27 days to decisionK231769 · Product code: **QLQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k231769/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Manual Instruments Designed For Use With Total Disc Replacement Devices (QLQ)
Date received	Jun 16, 2023
Decision date	Jul 13, 2023
Days to decision	27 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Aesculap Implants Systems, LLC</b>
Location	Center Valley, PA, US
Contact	Hamza Ahmed
510(k) history	3 submissions · 3 cleared · 2015-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k231769/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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