

K250399 Remanufactured EndoWrist Cadiere Forceps (420049)Aug 19, 2025
188 days to decisionK250399 · Product code: **QSM** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k250399/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Surgical, Computer Controlled Instrument, Remanufactured (QSM)
Date received	Feb 12, 2025
Decision date	Aug 19, 2025
Days to decision	188 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Rebotix
Location	St. Petersburg, FL, US
Contact	Chris Gibson
510(k) history	5 submissions · 5 cleared · 2024-2025

REGULATORY CONSULTANT

Consulting firm	Ajw Technology Consultants, Inc.
Contact	Alexsis Torres

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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