

**K250387 Remanufactured EndoWrist Long Tip Forceps  
(420048)**Aug 19, 2025  
189 days to decisionK250387 · Product code: **QSM** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k250387/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Surgical, Computer Controlled Instrument, Remanufactured (QSM)
Date received	Feb 11, 2025
Decision date	Aug 19, 2025
Days to decision	189 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

---

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

**REGULATORY CONSULTANT**

---

Consulting firm	<b>Ajw Technology Consultants, Inc.</b>
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)510k Database - [www.510kdatabase.net](https://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k250387/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](https://accessdata.fda.gov). - Generated May 14, 2026