

K234131 Resolv Endoscopic Hemostat SystemJul 26, 2024
211 days to decisionK234131 · Product code: **QAU** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k234131/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hemostatic Device For Endoscopic Gastrointestinal Use (QAU)
Date received	Dec 28, 2023
Decision date	Jul 26, 2024
Days to decision	211 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Hemostasis, LLC
Location	St. Paul, MN, US
Contact	Lakshmi Ganesh Bollina
510(k) history	10 submissions · 10 cleared · 2007-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k234131/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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