

K260287 SP Electrocautery Device (SP20)Feb 24, 2026
26 days to decisionK260287 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k260287/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 29, 2026
Decision date	Feb 24, 2026
Days to decision	26 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Single Pass, Inc.
Location	Lake Forest, CA, US
Contact	Greg Geissinger
510(k) history	2 submissions · 2 cleared · 2024-2026

REGULATORY CONSULTANT

Consulting firm	Geissinger Regulatory Consulting, Inc.
Contact	Greg Geissinger

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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