

**K232407 Cautery Protective Actuation System (CPAS)**Oct 6, 2023  
57 days to decisionK232407 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k232407/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 10, 2023
Decision date	Oct 6, 2023
Days to decision	57 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Safer Surgeries Medical Devices, LLC</b>
Location	Duluth, GA, US
Contact	Rowland Grattan
510(k) history	1 submissions · 1 cleared · 2023-2023

**REGULATORY CONSULTANT**

---

Consulting firm	<b>Secure Biomed Evaluations</b>
Contact	Braddon Linda

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/k232407/> 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 26, 2026