

**K183180 Caiman 5 Maryland**Mar 12, 2019  
113 days to decisionK183180 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k183180/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 19, 2018
Decision date	Mar 12, 2019
Days to decision	113 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Aesculap, Inc.</b>
Location	Burlingame, CA, US
Contact	Jessica Stigliano
510(k) history	207 submissions · 201 cleared · 1991-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k183180/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 28, 2026