

K223158 LigaSure™ XP Maryland Jaw Sealer/DividerJan 23, 2023
108 days to decisionK223158 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k223158/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 7, 2022
Decision date	Jan 23, 2023
Days to decision	108 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Covidien, LLC
Location	Mansfield, MA, US
Contact	Miranda Miles
510(k) history	88 submissions · 85 cleared · 2010-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k223158/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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