FDA 510(k) Premarket Notification

K223158 LigaSure™ XP Maryland Jaw Sealer/Divider

CLEARED

Jan 23, 2023
108 days to decision

K223158 · Product code: **GEI** · General & Plastic Surgery
Source: 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