FDA 510(k) Premarket Notification

K170107 Surgi Max Ultra

CLEARED

May 3, 2017
111 days to decision

K170107 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k170107/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 12, 2017
Decision date	May 3, 2017
Days to decision	111 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Elliquence, LLC**
Location	Shelton, CT, US
Contact	Paul D. Buhrke IV
510(k) history	8 submissions · 8 cleared · 2010-2025