FDA 510(k) Premarket Notification

CLEARED

K210942 MicroFrance Monopolar and Bipolar Electrosurgical Instruments

May 14, 2021
45 days to decision

K210942 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k210942/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 30, 2021
Decision date	May 14, 2021
Days to decision	45 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Integra Microfrance**
Location	Saint Aubin Le Monial Allier, FR
Contact	Malena Zammetti
510(k) history	1 submissions · 1 cleared · 2021-2021

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k210942/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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