FDA 510(k) Premarket Notification

K193611 Viveve 2.0 System

CLEARED

Jan 16, 2020
21 days to decision

K193611 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k193611/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 26, 2019
Decision date	Jan 16, 2020
Days to decision	21 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Viveve Medical, Inc.**
Location	Englewood, CO, US
Contact	Kevin Robison
510(k) history	3 submissions · 3 cleared · 2020-2021

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k193611/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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