FDA 510(k) Premarket Notification

K180584 Viveve RF System, Secure

CLEARED

Apr 6, 2018
32 days to decision

K180584 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k180584/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 5, 2018
Decision date	Apr 6, 2018
Days to decision	32 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Viveve, Inc.**
Location	Sunnyvale, CA, US
Contact	Suzon Lommel
510(k) history	3 submissions · 3 cleared · 2016-2022