FDA 510(k) Premarket Notification

K233002 BEED 2,5, BEED 3,8

CLEARED

Mar 20, 2024
180 days to decision

K233002 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k233002/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 22, 2023
Decision date	Mar 20, 2024
Days to decision	180 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Excelsior Resources, LLC**
Location	Ft. Lauderdale, FL, US
Contact	Christopher P Weber
510(k) history	1 submissions · 1 cleared · 2024-2024