FDA 510(k) Premarket Notification

K233374 Secret Pro

CLEARED

Dec 14, 2023
73 days to decision

K233374 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k233374/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 2, 2023
Decision date	Dec 14, 2023
Days to decision	73 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Ilooda Co,., Ltd.**
Location	Gwonseon-Gu, Suwon-Si, KR
Contact	Kathy Maynor
510(k) history	16 submissions · 16 cleared · 2015-2024