FDA 510(k) Premarket Notification

K220505 ATTIVA

CLEARED

Feb 9, 2023
352 days to decision

K220505 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k220505/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 22, 2022
Decision date	Feb 9, 2023
Days to decision	352 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

APPLICANT

Company	**Italian Engineering S.R.L.**
Location	Bologna, IT
Contact	Elisabetta Tonelli
510(k) history	1 submissions · 1 cleared · 2023-2023