FDA 510(k) Premarket Notification

K253915 MOVIVA® Hybrid Ablation Probe

CLEARED

Dec 18, 2025
10 days to decision

K253915 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k253915/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 8, 2025
Decision date	Dec 18, 2025
Days to decision	10 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Erbe Elektromedizin GmbH**
Location	Orange, CA, US
Contact	Matthias Kollek
510(k) history	15 submissions · 15 cleared · 1994-2026