

K232033 HYBRIDknife® flexNov 29, 2023
145 days to decisionK232033 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k232033/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 7, 2023
Decision date	Nov 29, 2023
Days to decision	145 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k232033/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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