

K230650 Monopolar Electrode Surgical InstrumentOct 12, 2023
217 days to decisionK230650 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k230650/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 9, 2023
Decision date	Oct 12, 2023
Days to decision	217 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Vhmed (Nantong) Co. , Ltd.
Location	Tongzhou Bay, CN
Contact	Guan Yunjuan
510(k) history	1 submissions · 1 cleared · 2023-2023

REGULATORY CONSULTANT

Consulting firm	P&L Scientific, Inc.
Contact	Jun Peng

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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