

K223222 Plasma IQJan 17, 2023
92 days to decisionK223222 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k223222/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 17, 2022
Decision date	Jan 17, 2023
Days to decision	92 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Neauvia North America, Inc.
Location	Raliegh, NC, US
Contact	Joy Willard
510(k) history	3 submissions · 3 cleared · 2020-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k223222/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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