

K230272 Renuvion® APR HandpieceApr 27, 2023
86 days to decisionK230272 · Product code: **GEI** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k230272/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 31, 2023
Decision date	Apr 27, 2023
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Apyx Medical Corporation
Location	Clearwater, FL, US
Contact	Angela Huber
510(k) history	5 submissions · 5 cleared · 2022-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k230272/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026