

**K231749 Radio Frequency System**Oct 19, 2023  
126 days to decisionK231749 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k231749/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 15, 2023
Decision date	Oct 19, 2023
Days to decision	126 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Shenzhensi Sincoheren S&amp;T Development Co., Ltd.</b>
Location	Shenzhen, CN
Contact	Cai Jialong
510(k) history	2 submissions · 2 cleared · 2023-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k231749/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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