

**K223239 8ch Wrist Coil**Dec 8, 2022  
49 days to decisionK223239 · Product code: **MOS** · Radiology  
Source: <https://www.510kdatabase.net/k223239/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Coil, Magnetic Resonance, Specialty (MOS)
Date received	Oct 20, 2022
Decision date	Dec 8, 2022
Days to decision	49 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Shenzhen RF Tech Co., Ltd.</b>
Location	Shenzhen, CN
Contact	Gary Wang
510(k) history	12 submissions · 12 cleared · 2015-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Bonnier Quality Supervision Consulting (Jm) Center</b>
Contact	Gary Wang

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k223239/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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