

**K223693 Electric breast pump (model: M5)**Mar 9, 2023  
90 days to decisionK223693 · Product code: **HGX** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k223693/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Breast, Powered (HGX)
Date received	Dec 9, 2022
Decision date	Mar 9, 2023
Days to decision	90 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Shenzhen Lutejiacheng Technology Co., Ltd.</b>
Location	Shenzhen, CN
Contact	Athena Pan
510(k) history	3 submissions · 3 cleared · 2022-2023

**REGULATORY CONSULTANT**

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Consulting firm	<b>Shenzhen Reanny Medical Devices Management Consulting Co., L</b>
Contact	Reanny Wang

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k223693/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 27, 2026