



April 7, 2026

Shenzhen Root Innovation Technology Co., Ltd.  
Caitlin Liang  
Official Correspondent  
#2-201, Floor 2 Hasee Computer Bldg.,  
# 2 Beier Rd., Bantian St., Longgang  
Shenzhen, 518129  
CHINA

Re: K253946

Trade/Device Name: Momcozy Wearable Breast Pump (S9 Pro, S9 Pro-A, S9 Pro-B,  
S9 Pro-C, S9 Pro-D, S9 ProY-A, S9 ProY-B, S9 ProY-C, S12 Pro,  
S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A,  
S12 ProY-B, S12 ProY-C)

Regulation Number: 21 CFR 884.5160

Regulation Name: Powered Breast Pump

Regulatory Class: II

Product Code: HGX

Received: March 6, 2026

Dear Caitlin Liang:

We have reviewed your section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (the Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. Although this letter refers to your product as a device, please be aware that some cleared products may instead be combination products. The 510(k) Premarket Notification Database available at <https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfpmn/pmn.cfm> identifies combination product submissions. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: The Center for Devices and Radiological Health (CDRH) does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, the Food and Drug Administration (FDA) may publish further announcements concerning your device in the Federal Register.

Additional information about changes that may require a new premarket notification are provided in the FDA guidance documents entitled "Deciding When to Submit a 510(k) for a Change to an Existing Device" (<https://www.fda.gov/media/99812/download>) and "Deciding When to Submit a 510(k) for a Software Change to an Existing Device" (<https://www.fda.gov/media/99785/download>).

Your device is also subject to, among other requirements, the Quality Management System Regulation (QMSR) (21 CFR Part 820), which includes, but is not limited to, ISO 13485 clause 7.3 (Design controls), ISO 13484 clause 8.3 (Nonconforming product), and ISO 13485 clause 8.5 (Corrective and preventative action). Please note that regardless of whether a change requires premarket review, the QMSR requires device manufacturers to review and approve changes to device design and production (ISO 13485 clause 7.3 and 21 CFR 820.70) and document changes and approvals in the device master record (21 CFR 820.181).

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR Part 803) for devices or postmarketing safety reporting (21 CFR Part 4, Subpart B) for combination products (see <https://www.fda.gov/combination-products/guidance-regulatory-information/postmarketing-safety-reporting-combination-products>); good manufacturing practice requirements as set forth in the Quality Management System Regulation (QMSR) (21 CFR Part 820) for devices or current good manufacturing practices (21 CFR Part 4, Subpart A) for combination products; and, if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR Parts 1000-1050.

All medical devices, including Class I and unclassified devices and combination product device constituent parts are required to be in compliance with the final Unique Device Identification System rule ("UDI Rule"). The UDI Rule requires, among other things, that a device bear a unique device identifier (UDI) on its label and package (21 CFR 801.20(a)) unless an exception or alternative applies (21 CFR 801.20(b)) and that the dates on the device label be formatted in accordance with 21 CFR 801.18. The UDI Rule (21 CFR 830.300(a) and 830.320(b)) also requires that certain information be submitted to the Global Unique Device Identification Database (GUDID) (21 CFR Part 830 Subpart E). For additional information on these requirements, please see the UDI System webpage at <https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory-assistance/unique-device-identification-system-udi-system>.

Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to <https://www.fda.gov/medical-devices/medical-device-safety/medical-device-reporting-mdr-how-report-medical-device-problems>.

For comprehensive regulatory information about medical devices and radiation-emitting products, including information about labeling regulations, please see Device Advice (<https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory-assistance>) and CDRH Learn (<https://www.fda.gov/training-and-continuing-education/cdrh-learn>). Additionally, you may contact the Division of Industry and Consumer Education (DICE) to ask a question about a specific regulatory topic. See the DICE website (<https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory->

[assistance/contact-us-division-industry-and-consumer-education-dice](#)) for more information or contact DICE by email ([DICE@fda.hhs.gov](mailto:DICE@fda.hhs.gov)) or phone (1-800-638-2041 or 301-796-7100).

Sincerely,

  
**Monica D. Garcia -S**

Monica D. Garcia, Ph.D.  
Assistant Director  
DHT3B: Division of Reproductive,  
Gynecology, and Urology Devices  
OHT3: Office of Gastrorenal, ObGyn,  
General Hospital, and Urology Devices  
Office of Product Evaluation and Quality  
Center for Devices and Radiological Health

Enclosure

## Indications for Use

510(k) Number (if known)  
K253946

Device Name

Momcozy Wearable Breast Pump (S9 Pro, S9 Pro-A, S9 Pro-B, S9 Pro-C, S9 Pro-D, S9 ProY-A, S9 ProY-B, S9 ProY-C, S12 Pro, S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C)

Indications for Use (Describe)

The Momcozy Wearable Breast Pump (S9 Pro, S9 Pro-A, S9 Pro-B, S9 Pro-C, S9 Pro-D, S9 ProY-A, S9 ProY-B, S9 ProY-C, S12 Pro, S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C) is a powered breast pump intended to express milk from lactating women in order to collect milk from their breasts. The device is intended for a single user.

Type of Use (Select one or both, as applicable)

Prescription Use (Part 21 CFR 801 Subpart D)

Over-The-Counter Use (21 CFR 801 Subpart C)

### CONTINUE ON A SEPARATE PAGE IF NEEDED.

This section applies only to requirements of the Paperwork Reduction Act of 1995.

**\*DO NOT SEND YOUR COMPLETED FORM TO THE PRA STAFF EMAIL ADDRESS BELOW.\***

The burden time for this collection of information is estimated to average 79 hours per response, including the time to review instructions, search existing data sources, gather and maintain the data needed and complete and review the collection of information. Send comments regarding this burden estimate or any other aspect of this information collection, including suggestions for reducing this burden, to:

Department of Health and Human Services  
Food and Drug Administration  
Office of Chief Information Officer  
Paperwork Reduction Act (PRA) Staff  
[PRASStaff@fda.hhs.gov](mailto:PRASStaff@fda.hhs.gov)

*"An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB number."*

## 510(k) Summary – K253946

### 1. Submitter Information

Applicant: Shenzhen Root Innovation Technology Co.,  
Ltd.  
Address: #2-201, Floor 2 Hasee Computer Building,  
No. 2 Beier Rd, Bantian Street, Longgang,  
Shenzhen 518129 China  
Tel.: 86-755-89698173

### 2. Correspondent Information

Contact: Ms. Caitlin Liang  
Shenzhen Root Innovation Technology Co., Ltd.  
Email: regulatory@momcozy.com

3. Date prepared: April 6, 2026

### 4. Device Information

Device Name: Momcozy Wearable Breast Pump (S9 Pro, S9 Pro-A, S9 Pro-B, S9 Pro-C,  
S9 Pro-D, S9 ProY-A, S9 ProY-B, S9 ProY-C, S12 Pro, S12 Pro-A, S12  
Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C)  
Common Name: Powered Breast Pump  
Regulation Number: 21 CFR 884.5160  
Regulation Name: Powered Breast Pump  
Product Code: HGX (Pump, Breast, Powered)  
Regulatory Class: Class II

### 5. Predicate Device Information

Device Name: Electric Breast Pump (Model LD-208L, LD-3010L, LD-2010L, LD-  
3010, LD-2010)  
510(k) Number: K241322  
Manufacturer: JOYTECH Healthcare Co.,Ltd.

The predicate device has not been subject to a design-related recall.

### 6. Device Description

The Momcozy Wearable Breast Pump (S9 Pro, S9 Pro-A, S9 Pro-B, S9 Pro-C, S9 Pro-D, S9 ProY-A, S9 ProY-B, S9 ProY-C, S12 Pro, S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C) are powered breast pumps intended to be used by lactating women to express and collect milk from their breasts; they are intended for a single user. The devices are powered by lithium battery, utilizing an embedded control program to manage all device functions. The main components of these pumps include: pump motor(main unit), milk collection set(include flange, diaphragm, silicone valve, milk collector, flange cover) and accessories( flange insert, charging cable). The devices use a diaphragm-type vacuum pump driven by a microprocessor. The user interface consists of buttons and LED display and allows the user to switch from stimulation, expression, and mixed modes (S12 Pro series only) and control the vacuum levels within those modes.

All available modes consist of 9 vacuum levels. The S9 Pro series is capable of providing vacuum levels from 65-290 mmHg with cycling rates from 52-141 cycles per minute in stimulation mode and vacuum levels from 130-290 mmHg with cycling rates from 18-74 cycles per minute in expression mode. The S12 Pro series is capable of providing vacuum levels from 80-160 mmHg with cycling rates from 73-142 cycles per minute in stimulation mode, vacuum levels from 130-290 mmHg with cycling rates from 18-44 cycles per minute in expression mode, and vacuum levels from 80-290 mmHg with cycling rates from 48-103 cycles per minute in mixed mode. The devices are charged with a 5 V DC adaptor and powered by an internal rechargeable lithium-ion polymer battery. The motor unit operates on embedded software. Software updates by end-users are not supported. The subject device is for repeated use by a single user in a home environment. The device is provided non-sterile.

The motor unit operates on a rechargeable battery and does not function when charging. The rechargeable battery can be charged from the external USB adapter if the motor unit is not in operation.

The breast pump expresses milk by creating a seal around the nipple using the flange and applying and releasing suction to the nipple. The milk is collected in a milk collection container, which can be used for storage. To prevent milk from flowing into the vacuum system, a backflow protection membrane physically separates the milk-contacting pathway from the vacuum system.

All other components (i.e., motor unit/housing) of the subject device is not in contact with the breast. All milk contacting components are compliant with 21 CFR 177.

## 7. Indications for Use

The Momcozy Wearable Breast Pump (S9 Pro, S9 Pro-A, S9 Pro-B, S9 Pro-C, S9 Pro-D, S9 ProY-A, S9 ProY-B, S9 ProY-C, S12 Pro, S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C) is a powered breast pump intended to express milk from lactating women in order to collect milk from their breasts. The device is intended for a single user.

## 8. Comparison of Intended Use and Technological Characteristics with the Predicate Device

The table below compares the intended use and technological characteristics of the subject and predicate device.

**Table 1: Comparator Table for Subject and Predicate Devices**

	<b>Momcozy Wearable Breast Pump (S9 Pro, S9 Pro-A, S9 Pro-B, S9 Pro-C, S9 Pro-D, S9 ProY-A, S9 ProY-B, S9 ProY-C, S12 Pro, S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C) K253946 Subject Device</b>	<b>Electric Breast Pump (Model LD-208L, LD-3010L, LD-2010L, LD-3010, LD-2010) K241322 Predicate Device</b>	<b>Comparison</b>
Code	HGX	HGX	Same
Indications for use	The Momcozy Wearable Breast Pump (S9 Pro, S9 Pro-A, S9 Pro-B, S9 Pro-C, S9 Pro-D, S9	The Electric Breast Pump (Model LD-208L, LD-3010L, LD-2010L, LD-3010, LD-2010) is a powered breast pump to be used by	Same

	ProY-A, S9 ProY-B, S9 ProY-C, S12 Pro, S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C) is a powered breast pump intended to express milk from lactating women in order to collect milk from their breasts. The device is intended for a single user.	lactating women to express and collect milk from their breasts. The Electric Breast Pump is intended for a single user.	
Single User	Yes	Yes	Same
Single/double pump	Single	Single or double	Different
Media separation (backflow protection)	Yes	Yes	Same
Cycling control mechanism	Microcontroller	Microcontroller	Same
Expression pattern	2-Phase	2-Phase	Same
Power supply	S9 Pro, S9 Pro-A, S9 Pro-B, S9 Pro-C, S9 Pro-D, S9 ProY-A, S9 ProY-B, S9 ProY-C: DC 3.7V/1600mAh Rechargeable lithium battery  S12 Pro, S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C: DC 3.7 V / 2*600mAh Rechargeable lithium battery	LD-3010L,LD-2010L: In:100 - 240 VAC 50/60 Hz 0.25A Mains Out: 5VDC, 1.0A Batt Out: 3.7VDC 2000mAh LD-3010,LD-2010: In:100 – 240 VAC 50/60 Hz 0.25A Mains Out: 5VDC, 1.0A Batt Out: 4 AA batteries	Different
Suction levels (expression)	S9 Pro, S9 Pro-A, S9 Pro-B, S9 Pro-C, S9 Pro-D, S9 ProY-A, S9 ProY-B, S9 ProY-C: 130-290 ± 20 mmHg  S12 Pro, S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C: 130 to -290(±20)mmHg	50-290 ± 30 mmHg	Different
Suction levels (stimulation)	S9 Pro, S9 Pro-A, S9 Pro-B, S9 Pro-C, S9 Pro-D, S9 ProY-A, S9 ProY-B, S9 ProY-C: 65-290 (± 20) mmHg  S12 Pro, S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C: 80 to -160(±20)mmHg	40-100 ± 30 mmHg	Different
Suction levels (mixed)	S12 Pro, S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C:	--	Different

	80 to -290(±20)mmHg		
Cycles per minute (expression)	S9 Pro, S9 Pro-A, S9 Pro-B, S9 Pro-C, S9 Pro-D, S9 ProY-A, S9 ProY-B, S9 ProY-C: 18 to 74 (±5)cycle/min S12 Pro, S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C: 18~44 (±5)cycle/min	25 to 65 (± 5) cycle/min	Different
Cycles per minute (stimulation)	S9 Pro, S9 Pro-A, S9 Pro-B, S9 Pro-C, S9 Pro-D, S9 ProY-A, S9 ProY-B, S9 ProY-C: 52 to 141(±5)cycle/min S12 Pro, S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C: 73 to 142(±5)cycle/min	80-120 ± 5 cpm	Different
Cycles per minute (mixed)	S12 Pro, S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C: 48 to 103 (±5)cycle/min	--	Differemt
Suction levels	9 levels stimulation, expression, mixed (S12 Pro series only)	9 expression, bionic, variable frequency, 6 stimulation	Different
Available modes	Stimulation, expression, mixed (S12 Pro series only)	Stimulate mode, Expression mode, Bionic sucking mode, Variable frequency pumping mode	Different
User Interface	On-Off switch, mode change button, vacuum adjustment buttons, LED	On-Off switch, mode change button, vacuum adjustment buttons, LED	Similar
Adjustable Suction Levels	Yes	Yes	Same
Mobile Application	No	No	Same
Design	Wearable	LD-3010L, LD-3010, LD-2010L, LD-2010: Tabletop LD-208L: Wearable	Same

The indications for use of the subject and predicate devices are identical, and both devices have the same intended use (i.e., for collection of breast milk from the breasts of lactating women).

The subject and predicate devices have similar technological features, including wearable operation, power supply, and user interface. However, as shown in the table above, there are technological differences between the subject and predicate devices, including different overall vacuum/cycle specifications, power supply, and available modes (mixed mode added to S12 Pro subject device). The different technological characteristics of the subject devices, as compared to the predicate device, do not raise different questions of safety and effectiveness.

## 9. Summary of Non-Clinical Performance Testing

### Biocompatibility

Biocompatibility information was provided in accordance with Attachment G of the 2023 FDA guidance document Use of International Standard ISO 10993-1, “Biological Evaluation of Medical Devices – Part 1: Evaluation and testing within a risk management process.”

### **Electrical Safety**

Testing was conducted in accordance with the following standards:

- ANSI/AAMI ES60601- 1:2005/A2:2010 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance),
- IEC 62133-2:2017, Secondary cells and batteries containing alkaline or other non-acid electrolytes - Safety requirements for portable sealed secondary cells, and for batteries made from them, for use in portable applications - Part 2: Lithium systems, and
- IEC 60601-1-11:2015 Medical electrical equipment – Part 1-11: General requirements for basic safety and essential performance – Collateral Standard: Requirements for medical electrical equipment and medical electrical systems used in the home healthcare environment.

### **Electromagnetic Compatibility**

Testing was conducted in accordance with the FDA Guidance “Electromagnetic Compatibility (EMC) of Medical Devices,” issued June 6, 2022 and IEC 60601-1-2:2014 Medical Electrical Equipment - Part 1-2: “*General Requirements For Basic Safety And Essential Performance - Collateral Standard: Electromagnetic Compatibility - Requirements And Tests.*”

### **Software**

Software was evaluated at the Basic Documentation level as recommended in the 2023 FDA guidance document “*Content of Premarket Submissions for Device Software Functions.*”

### **Performance Testing**

Other performance testing was conducted to show that the device meets its design requirements and performs as intended. The performance tests include:

- Vacuum level verification testing at each mode/cycle demonstrated that the devices meet mode/cycle specifications.
- Backflow protection testing was conducted to verify liquid does not backflow.
- Use life testing was conducted to demonstrate that the device maintains its specifications throughout its proposed use life.
- Battery performance testing was conducted to demonstrate that the battery remains functional during its stated battery use-life.
- Battery status indicator testing was conducted to demonstrate that the battery status indicator remains functional during its stated battery life.

## **10. Conclusion**

The results of the performance testing described above demonstrate that the Momcozy Wearable Breast Pump (S9 Pro, S9 Pro-A, S9 Pro-B, S9 Pro-C, S9 Pro-D, S9 ProY-A, S9 ProY-B, S9 ProY-C, S12 Pro, S12 Pro-A, S12 Pro-B, S12 Pro-C, S12 Pro-D, S12 ProY-A, S12 ProY-B, S12 ProY-C) is as safe and effective as the predicate device and support a determination of substantial equivalence.