July 21, 2022

STERIS Corporation
Gregory Land
Senior Regulatory Affairs Specialist
5960 Heisley Road
Mentor, Ohio 44060

Re: K213881
Trade/Device Name: Celerity Incubator
Regulation Number: 21 CFR 880.2800
Regulation Name: Sterilization process indicator
Regulatory Class: Class II
Product Code: FRC
Dated: July 5, 2022
Received: July 6, 2022

Dear Gregory Land:

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 (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 located 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: 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, FDA may publish further announcements concerning your device in the Federal Register.

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 803) for devices or postmarketing safety reporting (21 CFR 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 systems (QS) regulation (21 CFR Part 820) for devices or current good manufacturing practices (21 CFR 4, Subpart A) for combination products; and, if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 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) or phone (1-800-638-2041 or 301-796-7100).

Sincerely,

Clarence W. Murray, III, PhD
Assistant Director
DHT4B: Division of Infection Control
and Plastic Surgery Devices
OHT4: Office of Surgical
and Infection Control Devices
Office of Product Evaluation and Quality
Center for Devices and Radiological Health

Enclosure

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

Indications for Use

Form Approved: OMB No. 0910-0120
Expiration Date: 06/30/2020
See PRA Statement below.

510(k) Number *(if known)*
K213881

Device Name
Celerity Incubator

Indications for Use *(Describe)*
Use the Celerity Incubator to incubate and automatically read STERIS Celerity Biological Indicators for Steam and Vaporized Hydrogen Peroxide sterilization at 55°C-60°C for a fluorescent result.

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.

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510(k) Summary
For
Celerity Incubator

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Submission Date: July 21, 2022

K Number: K213881

1. Device Name

Trade Name:	**Celerity Incubator**
Device Class:	Class II
Common/usual Name:	Incubator
Classification Name:	Indicator, Biological Sterilization Process
Classification Number:	21 CFR 880.2800
Product Code:	FRC

2. Predicate Device

K190297 – Celerity HP Incubator

3. Reference Device

K173670 – Celerity Steam Incubator

4. Device Description

The Celerity Incubator incubates and reads fluorescent Vaporized Hydrogen Peroxide (VHP) and Steam Biological Indicators (BIs). The BI is incubated at 59°C with an acceptable tolerance of -4°C/+1°C. During incubation, the BI is monitored for a potential fluorescence signal generated as a result of the production of α-glucosidase. When a growth response has been detected or when the required incubation time has elapsed, the incubator indicates the results to the user.

4. Indications for Use

Use the Celerity Incubator to incubate and automatically read STERIS Celerity Biological Indicators for Steam and Vaporized Hydrogen Peroxide sterilization at 55°C - 60°C for a fluorescent result.

5. Technological Characteristics Comparison Table

A comparison of technical characteristics between the proposed and predicate devices is summarized in **Table 1**.

Table 1. Technological Characteristics Comparison Table to predicate device

Feature	Subject Device Universal Biological Indicator Incubator (K213881)	Predicate Device Celerity 20 HP Indicator (K190297)	Comparison
Indications for Use	Use the Celerity Incubator to incubate and automatically read STERIS Celerity Biological Indicators for Steam and Vaporized Hydrogen Peroxide sterilization at 55°C - 60°C for a fluorescent result.	Use the Celerity HP Incubator to incubate and automatically read Celerity 20 HP Biological Indicators at 57°C for a fluorescent result within 20 minutes.	Similar
Basis of Readout	Photodiode detects fluorescence produced by enzymatic activity that results from viable biological indicator organisms.	Photodiode detects fluorescence produced by enzymatic activity that results from viable biological indicator organisms.	Same
Incubation Temperature Range	55 - 60 °C	55 - 60 °C	Same
Voltage Range	100-240 VAC with 12 VDC conversion	100-240 VAC with 12 VDC conversion	Same
Test Capacity	8 wells	8 wells	Same
Calibration	Factory calibration – no calibration by customer	Factory calibration – no calibration by customer	Same
Incubation Time	20 minutes	20 minutes	Same
Fluorescence Detection	UV LEDs are used to excite the fluorescent molecule produced by enzyme cleavage of the fluorogenic substrate contained in the BI's media. The emitted light is detected by a photodiode.	UV LEDs are used to excite the fluorescent molecule produced by enzyme cleavage of the fluorogenic substrate contained in the SCBI's media. The emitted light is detected by a photodiode.	Same
Indication of Results	Positive – audible alarm, visual LED lights and screen Negative – optional alarm, visual indication with LED lights and LCD screen User must acknowledge results	Positive – audible alarm, visual LED lights and screen Negative – optional alarm, visual indication with LED lights and LCD screen User must acknowledge results	Same

Feature	Subject Device Universal Biological Indicator Incubator (K213881)	Predicate Device Celerity 20 HP Indicator (K190297)	Comparison
System Operation	The reader/incubator wells are arranged in 2 banks of 4 wells and preset to 59ºC. The user scans the barcode on the label of an activated BI using the system's barcode scanner and places it into an open well. The system detects the well the BI was placed into and begins measurement of fluorescence; a blinking yellow light indicates the incubation is in process and the read initiated. The System uses information from the barcode to apply the appropriate algorithm to the well. A "positive" reading is interpreted as an indication of a potential sterilization cycle failure. A "positive" finding is indicated to the user by red light on the front panel adjacent to the well, by an audible alarm, and by text displayed on the LCD screen. The alarm must be muted by the operator when a positive result is obtained. The LCD screen provides instructions for the user to turn off the alarm for that specific BI. Should another BI become "positive", the alarm will sound again and the above actions are repeated. If the result is not positive at the end of the incubation time, the result is negative. Negative results are identified by a green light on the front panel adjacent to the well with the "negative" BI and by text on the LCD. In addition, an optional alarm is available	The reader/incubator wells are arranged in 2 banks of 4 wells and preset to 57ºC. The measurement of fluorescence is initiated by placement of a Celerity HP BI into any of the incubation wells and pressing the adjacent "ACTION" button. When an SCBI is placed into a well, the auto-reader detects its presence. Upon pressing the button associated with that well, a blinking yellow light indicates that incubation is in process and the read initiated. A "positive" reading is interpreted as an indication of a potential sterilization cycle failure. A "positive" finding is indicated to the user by red light on the front panel adjacent to the well, by an audible alarm, and by text displayed on the LCD screen. The alarm must be muted by the operator when a positive result is obtained. The LCD screen provides instructions for the user to turn off the alarm for that specific BI. Should another BI become "positive", the alarm will sound again and the above actions are repeated. If the result is not positive at the end of the incubation time, the result is negative. Negative results are identified by a green light on the front panel adjacent to the well with the "negative" BI and by text on the LCD. In addition, an optional alarm is available	Similar

Feature	Subject Device Universal Biological Indicator Incubator (K213881)	Predicate Device Celerity 20 HP Indicator (K190297)	Comparison
	for negative results.	for negative results. User must acknowledge results.	

6. Summary of Non-Clinical Performance Testing

Testing was performed to evaluate performance and demonstrate that device met the acceptance criteria shown in **Table 2**.

Table 2. Performance Testing

Test	Acceptance Criteria	Result
Maintenance of Incubation Temperature	Temperature of BIs is maintained between 55°C – 60°C during incubation. Temperatures were observed over a 60min period and during a 1min loss of power.	PASS
Qualification Testing with intended Biological Indicators	Fluorescent Read meets ≥ 97% alignment with 7-day grow out per FDA guidance on reduced incubation time for intended Biological Indicators for both Steam and Vaporized Hydrogen Peroxide Celerity Biological Indicators	PASS
Human Factors	Typical users are capable of following the written instructions for use to correctly use the Celerity Incubators.	PASS
Electromagnetic Compatibility	IEC 60601-1-2:2014 General Requirements for Basic Safety and Essential Performance – Collateral Standard: Electromagnetic Compatibility – Requirements and Tests	PASS
Electrical Safety Conformance	IEC 60601-1 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	PASS
Software Validation	The software that controls the system was validated and determined to operate effectively and as designed.	PASS

7. Conclusion

The conclusions drawn from the non-clinical performance data demonstrates that the subject device is as safe, as effective, and performs as well as or better than the legally marketed predicate device (K190297), Class II (21 CFR 880.2800), product code FRC.