



July 29, 2026

CenterPoint Systems, LLC
Lauren Marold
Senior Regulatory Specialist
3338 Pkwy. Blvd.
West Valley City, Utah 84119

Re: K261766

Trade/Device Name: CPS Locator 3D Plus Delivery Catheter
Regulation Number: 21 CFR 870.1250
Regulation Name: Percutaneous Catheter
Regulatory Class: Class II
Product Code: DQY
Dated: June 29, 2026
Received: June 30, 2026

Dear Lauren Marold:

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: 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.

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 13485 clause 8.3 (Nonconforming product), ISO 13485 clause 8.5.2 (Corrective action), and ISO 13485 clause 8.5.3 (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 ISO 13485 clause 7.5) and document changes and approvals in the Medical Device File (ISO 13485 clause 4.2.3).

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) or phone (1-800-638-2041 or 301-796-7100).

Sincerely,

JESSICA L.
BATISTA-S

Digitally signed by JESSICA
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Date: 2026.07.29 13:03:51
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Jessica Batista Bertolini
Acting Assistant Director
Division of Cardiac Electrophysiology,
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Center for Devices and Radiological Health

Enclosure

Indications for Use

510(k) Number (if known)

K261766

Device Name

CPS Locator 3D Plus Delivery Catheter

Indications for Use (Describe)

The Delivery Catheter is indicated for the introduction of various types of pacing or defibrillator leads to the right heart, inclusive of the Left Bundle Branch Area.

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.

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510(k) Summary

1.1 Submitter

Name CenterPoint Systems LLC

Address 3338 Parkway Blvd
West Valley City UT

Phone 860-707-3892

Contact Person: Lauren Marold, Senior Regulatory Specialist

Date Prepared: 11 June 2026

1.2 Device

Name of Device: CPS Locator 3D Plus Delivery Catheter

Classification Name: Percutaneous Catheter

Regulatory Class: Class II per 21 CFR 870.1250

Product Code: DQY

1.3 Predicate Device

Predicate Name and 510(k) Number: CPS Locator 3D Plus Delivery Catheter, K250828

This predicate has not been subject to a design-related recall. The reference device, CPS Locator 3D Delivery Catheter, K250147, has not been subject to a design-related recall.

1.4 Device Description

The modified Delivery Catheter (CPS Locator 3D Delivery Catheter) is a single-use percutaneous catheter indicated for the introduction of various types of pacing or defibrillation leads to the right heart, inclusive of the Left Bundle Branch Area.

The modified Delivery Catheter is packaged with a dilator for introduction into the vasculature. Proximally, the modified Delivery Catheter is equipped with a hemostatic valve, and the distal soft, rounded, radiopaque tip facilitates imaging under fluoroscopy. The modified Delivery Catheter is designed to be slittable, thereby allowing its removal after device placement. A variety of curves and lengths are available to accommodate various anatomies and different locations.

1.5 Indications for Use

The Delivery Catheter is indicated for the introduction of various types of pacing or defibrillation leads to the right heart, inclusive of the Left Bundle Branch Area.

1.6 Comparison of Technological Characteristics with the Predicate Devices

The Proposed Device and Predicate Device are similar in indications for use, intended use, technological characteristics, and principles of operation. The Proposed Device and the Reference Device have the same indications for use and intended use, and are similar in technological characteristics and principles of operation.

The differences between the Proposed Device and the Predicate Device are minor and raise no different questions of safety and effectiveness. In accordance with 21CFR807.92(a)(6) a summary of how the technological characteristics of the Proposed Device compares to the Predicate Device and Reference Device is provided below.

Feature	CPS Locator 3D Plus Delivery Catheter (subject device)	Primary Predicate: CPS Locator 3D Plus Delivery Catheter (K250828)	Reference Device: CPS Locator 3D Delivery Catheter (K250147)	Same / Different between Proposed & Predicates
Intended Use	Percutaneous catheter for the delivery of leads	Percutaneous catheter for the delivery of leads	Percutaneous catheter for the delivery of leads	Same
Indications for Use	The Delivery Catheter is indicated for the introduction of various types of pacing or defibrillator leads to the right heart, inclusive of the Left Bundle Branch Area.	The Delivery Catheter is indicated for the introduction of various types of catheters and pacing or defibrillator leads.	The Delivery Catheter is indicated for the introduction of various types of pacing or defibrillator leads to the right heart, inclusive of the Left Bundle Branch Area.	Substantially Equivalent to Predicate Device Same as Reference Device
Product Code	DQY	DQY	DQY	Same
Regulation number	21 CFR 870.1250	21 CFR 870.1250	21 CFR 870.1250	Same
Prescription Device	Yes	Yes	Yes	Same
Catheter Type	Percutaneous Catheter	Percutaneous Catheter	Percutaneous Catheter	Same
Guidewire Compatibility	0.035"	0.035"	0.035"	Same
Outer Diameter	10.3F	10.2F	9.0F	Substantially Equivalent

Feature	CPS Locator 3D Plus Delivery Catheter (subject device)	Primary Predicate: CPS Locator 3D Plus Delivery Catheter (K250828)	Reference Device: CPS Locator 3D Delivery Catheter (K250147)	Same / Different between Proposed & Predicates
Inner Diameter	8.5F	8.4F	7.0F	Substantially Equivalent
Working Length	42cm, 45cm, 48cm	42cm, 45cm	42cm, 45cm	Substantially Equivalent
Components Provided	Catheter, Dilator, Luer cap	Catheter, Dilator, Luer cap	Catheter, Dilator, Luer cap	Same
Hydrophilic Liner	Yes	Yes	Yes	Same
Radiopaque Distal Tip	Yes	Yes	Yes	Same
Valve	Yes	Yes	Yes	Substantially Equivalent
Shaft Materials	PEBAX with Barium Sulfate and colorants	PEBAX with Barium Sulfate and colorants	PEBAX with Barium Sulfate and colorants	Same
Braid Reinforcement	Yes	Yes	Yes	Same
Dilator	Yes	Yes	Yes	Same
Multiple Distal End Shapes Available	Yes	Yes	Yes	Same
Sterility	E-Beam Sterilization, SAL 10^{-6}	E-Beam Sterilization, SAL 10^{-6}	E-Beam Sterilization, SAL 10^{-6}	Same
Number of Uses	Single patient use	Single patient use	Single patient use	Same

Feature	CPS Locator 3D Plus Delivery Catheter (subject device)	Primary Predicate: CPS Locator 3D Plus Delivery Catheter (K250828)	Reference Device: CPS Locator 3D Delivery Catheter (K250147)	Same / Different between Proposed & Predicates
Principles of Operation	After venous access is gained, the catheter and dilator are advanced over a guidewire to the desired location. The dilator is removed and a catheter or lead is placed through the Delivery Catheter. The Delivery Catheter may be removed by slitting.	After venous access is gained, the catheter and dilator are advanced over a guidewire to the desired location. The dilator is removed and a catheter or lead is placed through the Delivery Catheter. The Delivery Catheter may be removed by slitting.	After venous access is gained, the catheter and dilator are advanced over a guidewire to the desired location. The dilator is removed and a catheter or lead is placed through the Delivery Catheter. The Delivery Catheter may be removed by slitting.	Same

The modified Delivery Catheter is used for the same intended use in the same anatomical location using the same principles of operation as the predicate device.

1.7 Performance Data

All necessary performance testing has been conducted on the CPS Locator 3D Plus Delivery Catheter to assure substantial equivalence to the predicate devices and to demonstrate the device performs as intended. All testing was performed on test units representative of finished devices.

The device passed the following tests, which were conducted in accordance with noted standards:

- Sterilization validation per ISO 11137
- Packaging validation per ANSI/AAMI/ISO 11607-1
- Visual Inspection
- Flush test
- Tensile tests
- Simulated use testing, including use/compatibility with ancillary devices
- Valve liquid leak test
- Sheath and Dilator Dimensional verification, including OD/ID, working length

1.8 Conclusions

Based on the similarity of the subject and predicate devices in terms of the intended use, principle of operation and overall technological characteristics, and the results of the testing summarized in section 1.7, the CPS Locator 3D Plus Delivery Catheter is substantially equivalent to the predicate device (K250828).