

AUG 12 2003

Endocare, Inc.
Traditional 510(k): Cryocare CS Surgical System

K 032333



510(K) SUMMARY

Prepared May 7, 2003

| | | | |
|--------------------|--|---------|---|
| TRADE NAME | Cryocare CS Surgical System | | |
| COMMON NAME | Cryosurgical unit and accessories | | |
| CLASSIFICATION | Class II (21 CFR 878.4350) | | |
| SUBMITTED BY | Endocare, Inc. 201 Technology Irvine, CA 92618 | CONTACT | Eben Gordon Regulatory Affairs 949.450.5424 949.450.5300 |
| PREDICATE DEVICE | K023757 - Endocare Cryocare Surgical System Decision date: 12/5/2002 K002615 - Endocare Cryocare Surgical System with CryoGuide Decision date: 11/13/2000 K030191 - Terason Model 2000/BAS Portable Ultrasound System Decision date: 1/29/2003 | | |
| DEVICE DESCRIPTION | <p>The Cryocare CS (Combo System) Surgical System consists of a compact, easy-to-operate console that delivers cold temperatures to targeted tissue (via connected CryoProbes) and monitors temperatures in the surrounding tissue (via connected TempProbes).</p> <p>The Cryocare CS Surgical System has a fold-down 19" LCD high-resolution display screen, a video printer for hard copy prints of the captured images and patient information, a CD-R/W drive for data storage and retrieval, an alphanumeric keypad and a remote keypad.</p> <p>The Cryocare CS unit integrates the functions of the existing Cryocare surgical system and the CryoGuide system into one console. The CryoGuide software is an optional brachy-like intraoperative real-time guidance and planning system, which identifies and guides CryoProbe placement for prostate procedures.</p> <p>An added feature to the Cryocare CS Surgical System is integration of the Terason 2000 ultrasound trans-rectal probe for dedicated prostate imaging. The ultrasound module utilizes the Terason Model 2000 Handheld Ultrasound System using Model 8EC4 Endocavity Smart Probe. It is capable of transverse and sagittal views, and operates in B, Power Doppler, Color Doppler (including directional and non-directional Power Doppler), and Harmonic Imaging. This use includes ultrasound guidance for placement of needles, cryosurgery, and brachytherapy.</p> <p>Ultrasound images may be obtained either from an external ultrasound (via video connection) or from the integrated Terason 2000 ultrasound module. The user screen simultaneously displays the real-time ultrasound image along with the treatment screen. For prostate procedures, the treatment screen displays the prostagraph showing the anatomical CryoProbe and TempProbe placements and</p> | | |

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associated temperature readings.

The Cryocare CS Surgical System can control up to eight, single-use, disposable CryoProbes and monitor up to eight independent TempProbes. The console operates off standard 120/230 VAC (60/50 Hz) wall power and utilizes inert argon and helium gas. An IBM-compatible microprocessor serves as the host computer operating in a Windows environment. CryoProbe control can also be achieved via the remote control keypad. The CryoProbes can be operated manually or using the AutoFreeze mode, which allows users to pre-program specific prostate treatment parameters.

**INDICATIONS FOR
USE**

The Cryocare CS Surgical System has the **same intended use** as previously cleared for the Cryocare CS Surgical System - K023757.

The Cryocare CS Surgical System is intended for use in open, minimally invasive or endoscopic surgical procedures in the areas in general surgery, urology, gynecology, oncology, neurology, dermatology, ENT, proctology, pulmonary surgery and thoracic surgery. The system is designed to freeze/ablate tissue by the application of extreme cold temperatures including prostate and kidney tissue, liver metastases, tumors, skin lesions, and warts.

The intended uses of the Terason 2000 Ultrasound System, when used as a part of the Cryocare CS Surgical System, are a subset of the uses cleared under K030191, and are documented in section 4.3 of this submission.

**SUMMARY OF
SUBSTANTIAL
EQUIVALENCE**

The Cryocare CS Surgical System has the following similarities to that of the previously cleared predicate devices:

- Has the same intended use,
- Use the same operating principle and has not altered the fundamental technology,
- Incorporate the same CryoProbe,
- Incorporate the same system design,
- Incorporate the same patient contacting materials,
- Same manufacturing materials, and
- Packaged and sterilized using the same materials and processes.

In summary, the modified Cryocare CS Surgical System described in this submission is, in our opinion, substantially equivalent to the predicate devices.



FEB 21 2008

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Endocare, Inc.
c/o Ms. Laura Danielson
510(k) Program Manager
TUV Product Service
1775 Old Highway 8 NW
New Brighton, MN 55112-1891

Re: K032333
Trade Name: Cryocare CS Surgical System
Regulation Number: 21 CFR 878.4350
Regulation Name: Cryosurgical unit and accessories
Regulatory Class: II (two)
Product Code: OCL, GEH, IYN, IYO, ITX
Dated: July 28, 2003
Received: July 29, 2003

Dear Ms. Danielson:

This letter corrects our substantially equivalent letter for this application.

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 (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. 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.

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.

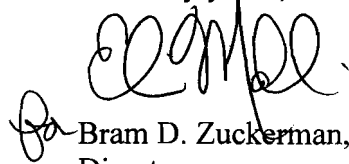
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Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other 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); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to continue marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at (240) 276-0120. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (240) 276-3150 or at its Internet address <http://www.fda.gov/cdrh/dsma/dsmamain.html>

Sincerely yours,



Bram D. Zuckerman, M.D.
Director
Division of Cardiovascular Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

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Indications for Use Statement

510(k) Number: K 032333

Device Name: Cryocare CS Surgical System

Indications for Use:

The Cryocare CS Surgical System is intended for use in open, minimally invasive or endoscopic surgical procedures in the areas in general surgery, urology, gynecology, oncology, neurology, dermatology, ENT, proctology, pulmonary surgery and thoracic surgery. The system is designed to freeze/ablate tissue by the application of extreme cold temperatures including prostate and kidney tissue, liver metastases, tumors, skin lesions, and warts.

In addition, the system is intended for use in the following indications:

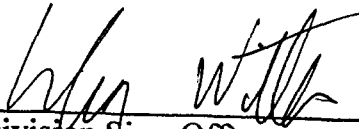
General Surgery

- Destruction of warts or lesions
- Palliation of tumors of the oral cavity, rectum and skin
- Ablation of leukoplakia of the mouth, angiomas, sebaceous hyperplasia, basal cell tumors of the eyelid or canthus area, ulcerated basal cell tumors, dermatofibromas, small hemanglomas, mucocele cysts, multiple warts, plantar warts, hemorrhoids, anal fissures, perianal condylomata, pilonidal cysts, actinic and seborrheic keratoses, cavernous hemanglomas, recurrent cancerous lesions

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use X (Per 21 CFR 801.109)



(Division Sign-Off)
Division of General, Restorative
and Neurological Devices

510(k) Number 032333

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Indications for Use Statement (Continued)

Urology

- Ablation of prostate tissue in cases of prostate cancer and benign prostatic hyperplasia

Gynecology

- Ablation of malignant neoplasia or benign dysplasia of the female genitalia

Oncology

- Ablation of cancerous or malignant tissue
- Ablation of benign tumors
- Palliative intervention

Neurology

- Freezing of nerve tissue in pain management/cryoanalgesia

Dermatology

- Ablation or freezing of skin cancers and other cutaneous disorders

Proctology

- Ablation of benign or malignant growths of the anus or rectum
- Ablation of hemorrhoids

Thoracic Surgery

- Ablation of arrhythmic cardiac tissue
- Ablation of cancerous lesions

**PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE
IF NEEDED)**

Concurrence of CDRH, Office of Device Evaluation (ODE)

PRESCRIPTION USE X (PER 21 CFR 801.109)