

Tel 513-772-8810 Fax 513-772-9119 www.genthermcsz.com

## **URGENT FIELD SAFETY NOTICE**

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Blanketrol II Hyper-Hypothermia System FA2019-004,010
Correction

8/1/2019

This is to inform you that Gentherm Medical, LLC (formerly Cincinnati Sub-Zero) is conducting a voluntary product recall.

For Attention of: Blanketrol II user

**Details on affected devices:** Blanketrol II Hyper-Hypothermia System, Models 222R and 222S, Parts 86006 (222R, 100V), 86106 (222R, 115V), 86109 (222R, 240V), 86165 (222S, 115V), 86206 (222R, 230V), 86265 (222S, 230V)

Affected Serial Numbers: Model 222R (071-18490 - 123-20534)

Model 222S (104-2-00001 - 161-2-01848)

**Description of device:** The BLANKETROL II Hyper-Hypothermia System is used to lower or to raise a patient's temperature and/or maintain a desired patient temperature through conductive heat transfer.

See enclosed product labels in Appendix A for ease in identifying the product at the user level.

**Description of the problem:** The purpose of this letter is to advise you that warnings have been clarified stating that exceeding 40°C for extended periods may cause tissue damage. See Appendix B for specific changes that have been made to the device manuals.

Immediately examine your inventory and update manual(s) subject to recall. In addition, if you may have further distributed this product, please identify your customers and notify them at once of this product recall. Your notification to your customers may be enhanced by including a copy of this correction letter.

#### **INSTRUCTIONS TO CUSTOMERS:**

- Access updated manuals and ensure that obsolete manuals are removed from service. Updated manuals may be accessed via <a href="www.gentherm.com">www.gentherm.com</a> or physical copies may be requested from Gentherm Medical, LLC at 1-888-437-5608.
- 2) Ensure that all users are informed of the contents of this letter. If you have further distributed this product, please provide those accounts with a copy of this notice.
- 3) Please complete and return the enclosed response form as soon as possible to acknowledge receipt of this notification and to inform Gentherm Medical, LLC that you have performed and completed the requested actions. Return the form by e-mail to <u>FA2019</u>-004 010@gentherm.com or mail to:

Gentherm Medical, LLC 12011 Mosteller Road Cincinnati, OH 45241





Tel 513-772-8810 Fax 513-772-9119 www.genthermcsz.com

The Competent (Regulatory) Authority of your country has been informed about this communication to customers.

### Transmission of this Field Safety Notice:

This notice needs to be passed on to all those who need to be aware within your organization or to any organization where the potentially affected devices have been transferred. (As appropriate)

Please transfer this notice to other organizations on which this action has an impact. (As appropriate)

Please maintain awareness on this notice and resulting action for an appropriate period to ensure effectiveness of the corrective action.

Please report all device-related incidents to the manufacturer, distributor or local representative, and the national Competent Authority if appropriate, as this provides important feedback.

### Contact reference person:

Stephanie Vocke Gentherm Medical, LLC 12011 Mosteller Road (513)719-3262

The undersigned confirms that this notice has been provided to the appropriate Regulatory Agency.





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## **Appendix A: Blanketrol II Product Labels**



DANGER: RISK OF EXPLOSION DO NOT USE IN THE PRESENCE OF FLAMMABLE ANESTHETICS.

SN

**222R** 

11.6A 100V~ 50/60Hz

R134a



DANGER: RISK OF ELECTRICAL SHOCK, DISCONNECT POWER BEFORE SERVICING.





Cincinnati Sub-Zero Products, Inc. 12011 Mosteller, Chompati, Orb., U.S.A. 45241 (513) 772-8310 e-mail: eszne@eszinc.com

56629-J

# BLANKETROL® II

**CAUTION:** FEDERAL (U.S.A.) LAW RESTRICTS THIS DEVICE TO SALE BY OR ON THE ORDER OF A PHYSICIAN.

DANGER: RISK OF EXPLOSION DO NOT USE IN THE PRESENCE OF FLAMMABLE ANESTHETICS.

SN

222R

10.2A 115V~ 50/60Hz

REFRIGERANT R134a 15oz.

<u>DESIGN PRESSURES</u> HIGH SIDE 240 PSIG LOW SIDE 240 PSIG



DANGER: RISK OF ELECTRICAL SHOCK, DISCONNECT POWER BEFORE SERVICING.





Cincinnati Sub-Zero Products, Inc. 12011 Mosteller, Cincinnati, Ohio, U.S.A. 45241 (513) 772-8810 9-mail: cszinc@cszinc.com

56628-L



58684-N

Tel 513-772-8810 Fax 513-772-9119 www.genthermcsz.com

# 222R - 50/60Hz SN TEMP. CLASS N R134a (15 oz.) Cincinnati Sub-Zero C€<sub>0344</sub>

Cincinnati Sub-Zero Products, Inc. 12011 Mosteller, Cincinnati, Ohio, U.S.A. 45241 (513) 772-8810

e-mail: cszinc@cszinc.com

222R 50/60Hz SN TEMP. CLASS N R134a (15 oz.) Cincinnati Sub-Zero Cincinnati Sub-Zero Products, Inc. 12011 Mosteller, Cincinnati, Chio, U.S.A. 45241 (513) 772-8819 56681-N



Tel 513-772-8810 Fax 513-772-9119 www.genthermcsz.com

CAUTION: FEDERAL (U.S.A.) LAW RESTRICTS THIS DEVICE TO SALE BY OR ON THE ORDER OF A PHYSICIAN.

**DANGER: RISK OF EXPLOSION** DO NOT USE IN THE PRESENCE OF FLAMMABLE ANESTHETICS.

SN

**REF 222S** 50/60Hz

> REFRIGERANT R134a 15oz.

**DESIGN PRESSURES** HIGH SIDE **LOW SIDE** 

**240 PSIG 240 PSIG** 





MODEL 2228 MEDICAL **ELECTRICAL EQUIPMENT** WITH RESPECT TO ELECTRIC SHOCK, FIRE AND MECHANICAL HAZARDS ONLY IN ACCORDANCE WITH UL60601-1, IEC60601-1, IEC60601-2-35, ASTM F2196-02, IEC60601-1-2 AND CAN/CSA-C22.2 No. 601.1



Cincinnati Sub-Zero Products, Inc. 12011 Mosteller, Cincinnati, Ohio, U.S.A. 45241 (513) 772-8810 e-mail: cszinc@cszinc.com

DANGER: RISK OF ELECTRICAL SHOCK, DISCONNECT POWER BEFORÉ SERVICING.

**C**€ 0344

57037-C

**DANGER: RISK OF EXPLOSION** DO NOT USE IN THE PRESENCE OF FLAMMABLE ANESTHETICS.

**DANGER: RISK OF ELECTRICAL** SHOCK, DISCONNECT POWER BEFORE SERVICING.

SN

**REF 222S** 

50/60Hz

REFRIGERANT R134a 15oz.

**DESIGN PRESSURES** HIGH SIDE **240 PSIG LOW SIDE 240 PSIG** 





MODEL 222S MEDICAL ELECTRICAL EQUIPMENT WITH RESPECT TO ELECTRIC SHOCK, FIREAND MECHANICAL HAZARDS ONLY IN ACCORDANCE WITH UL80801-1, EC80801-1, IEC60801-2-35, ASTM F2196-02, IEC60601-1-2, AND CAN/CSA-C22.2 No. 601.1



Cincinnati Sub-Zero

Cincinnati Sub-Zero Products, Inc. 12011 Mosteller, Cincinnati, Ohio, U.S.A. 45241 (513) 772-8810 website: cszmedical.com e-mail: cszinc@cszinc.com

C € 0344

57038-B



Tel 513-772-8810 Fax 513-772-9119 www.genthermcsz.com

# Appendix B: Blanketrol III Hyper-Hypothermia System Manual Updates

Affected Manual	Updates		
nanges are designated in RED se updated manual(s) for more details.			
56230 (222R Operation and Technical Manual): WARNINGS section  57126 (222R Operation Manual): WARNINGS section	WARNING: A physician's order is required for setting blanket temperature and use of equipment. At least every 20 minutes, or as directed by physician, check patient's temperature and skin condition of areas in contact with blanket; also, check blanket water temperature. Pediatric, temperature-sensitive, patients with vascular disease and operating room patients are at greater risk for developing tissue injuries, and this should be considered when selecting the temperature, duration of therapy and frequency of skin checks. Notify the physician promptly of any change in patient status in order to avoid serious injury or death.  WARNING: The method of temperature control provided by all hyperhypothermia units presents the danger of heating or cooling body tissues, particularly the skin, to a point where they are injured, i.e., burns or frostbite, respectively. The clinician is responsible for determining the appropriateness of the temperature limits in dependency to time. Exceeding 40°C water temperature for extended periods can cause tissue damage and burns. Clinical judgment should be used to determine the safe maximum contact periods based on patient age, clinical condition and current medications. Depending on the extent and severity of a burn, very serious and even fatal complications may arise.		
57947 (222S Operation and Technical Manual): WARNINGS section, Sections 2-4, 3-2, 3-3, and 3-4 56277 (222S Operation Manual): WARNINGS section, Sections 3-2, 3-3 and 3-4	WARNING: A physician's order is required for setting blanket temperature and use of equipment. At least every 20 minutes, or as directed by the physician, check patient's temperature and skin integrity of areas in contact with blanket; also, check the BLANKETROL II's water temperature. Pediatric patients, temperature-sensitive patients with vascular disease, surgical patients, diabetics and patients with Raynaud's Disease at greater risk for developing tissue injuries, and this should be considered when selecting the temperature, duration of therapy and frequency of skin checks. Notify the physician promptly of any change in patient status in order to avoid serious injury or death.		
57947 (222S Operation and Technical Manual): WARNINGS section and Section 2-4 56277 (222S Operation Manual): WARNINGS section	WARNING: The method of temperature control provided by all hyper-hypothermia units presents the danger of heating or cooling body tissues, particularly the skin, to a point where they are injured, i.e. burns or frostbite, respectively. The clinician is responsible for determining the appropriateness of the temperature limits in dependency to time. Exceeding 40°C water temperature for extended periods can cause tissue damage and burns. Clinical judgment should be used to determine the safe maximum contact periods based on patient age, clinical condition, and current medications. Depending on the extent and severity of a burn, very serious and even fatal complications may arise.		





Please check ALL appropriate boxes.

Corporate Office 12011 Mosteller Road Cincinnati, Ohio 45241-1528

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## **URGENT FIELD SAFETY NOTICE**

## **Response Form**

Please complete this form after your facility has performed the instructions provided in the correction letter. Return the completed form by mail to Gentherm Medical, LLC, 12011 Mosteller Road, Cincinnati, OH 45241, by fax to (513)772-9119 or scan and e-mail to FA2019-004 010@gentherm.com.

I have read and understand the field notif		
I have ensured all users are informed of	the contents of this letter.	
Indicate disposition of recalled product:		
☐ Corrected:		
(Specify serial number	er(s) and date)	
☐ Returned:		
(Specify serial numbe	er(s), and date)	
☐ Destroyed:		
(Specify serial number		
I have identified and notified my custome	ers that were shipped or may have been shipped this	product by;
Attached is a list of customers who receive to notify my customers.	OR ved/may have received this product. I would like Ger	ntherm <b>Medic</b> al
Signature	Date	
Printed Name	Email Address	
Facility Name	Facility Address, City, State, Zip Code	
Phone Number		

