600 N. Field Drive
Lake Forest, IL 60045
www.icumed.com

22 October 2024

URGENT MEDICAL DEVICE SAFETY ALERT

Counterfeit Batteries Used with Plum 360™, Plum A+™ & Plum A+3™ Infusion Systems

Product Name	Pump List Number
Plum 360™ Infusion System	30010
Plum A+™ & Plum A+3™ Infusion Systems	11005, 11971, 12391, 12618, 20678, 20679, 20792, 60529, 12348, 11973

Dear Valued Plum Infusion System Customers:
Director of Biomedical Engineering
Director of Nursing
Director of Risk Management

Issue:
ICU Medical has received reports of counterfeit CSB batteries being used with Plum Infusion Systems. While these counterfeit batteries are visually similar to the Plum batteries supplied by ICU Medical, they are in fact not the same batteries and have not been tested or validated for use with Plum Infusion Systems. Preliminary reports suggest that these counterfeit batteries fail to hold their charge and the pump may display messages to replace batteries earlier than expected. OSI Batteries and their customers are distributing these counterfeit CSB batteries without authorization from ICU Medical.

Do not use spare parts that are not authorized by ICU Medical. Please ensure that any entity providing service or repair activities for your Plum Infusion Systems uses only parts authorized or directly provided by ICU Medical.

Specific to these counterfeit batteries, please use the following information to identify counterfeit batteries.

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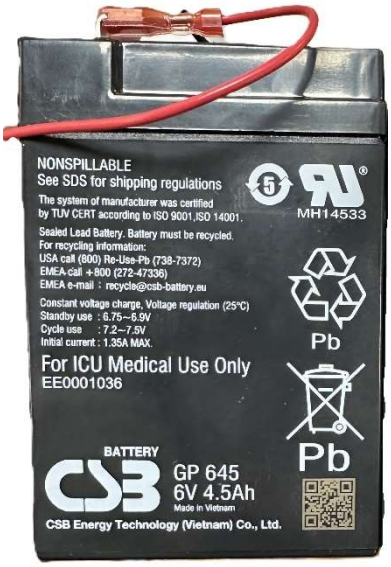
AUTHORIZED CSB BATTERY ACCEPTABLE TO USE	COUNTERFEIT CSB BATTERY DO NOT USE
	
DESCRIPTION	
Battery produced by CSB for ICU Medical	Counterfeit battery that was not produced for ICU Medical or tested to ICU Medical specifications
LABELING	
□ **WHITE TEXT** □ ICU Medical Test Label (#1) □ CE Mark (#2) □ Includes "For ICU Medical Use Only" (#3) □ Date code (yellow label found on side of battery): 2404xxxx and higher **OR** C2404XXX and higher	□ **WHITE TEXT** □ **NO** ICU Medical Test Label □ **NO** CE Mark □ Includes "For ICU Medical Use Only" □ Date code (yellow label found on side of battery): W2401xxxx - W2406xxx
ICU PART NUMBER ON THE BOX	
SUB0000864	N/A

Potential Risk:

Preliminary information indicates that counterfeit batteries may have a substantially diminished life.

If the pump is running on battery power and the pump has triggered the replace battery alarm, the user may not have sufficient time to plug the pump into AC power after the Low Battery alarm is activated, which may result in an interruption or delay of therapy. An interruption or delay of therapy can lead to serious patient injury or death, depending on the clinical situation and the type of medication being administered. **To date, ICU Medical has not received any adverse event reports potentially related to this issue.**

Actions to be taken by the Customer:

There is no need to return or discontinue using your Plum 360, Plum A+, or Plum A+3 pumps.

1. Identify if you have counterfeit batteries using the guidance provided above.
2. Remove counterfeit batteries from use and inventory and destroy them per hospital guidelines.

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Actions for Biomedical Engineering:

1. Ensure all users or potential users of these pumps are immediately made aware of this alert.
2. Complete and return the attached Response Form to icumedical7561@sedgwick.com **within ten days of receipt** to acknowledge your understanding of this alert.
3. **DISTRIBUTORS:** If you have distributed potentially affected products to your customers, please immediately forward this alert to them. Request that they complete the response form and return it to icumedical7561@sedgwick.com.

Actions by ICU Medical:

ICU Medical is providing this communication to inform you of counterfeit batteries that should not be used with the Plum Infusion Systems.

For further inquiries, please contact ICU Medical using the information provided below.

ICU Medical Contact	Contact Information	Areas of Support
Global Complaint Management	1-844-654-7780 (M-F, 8:00am – 5:00pm CT) or ProductComplaintsPP@icumed.com	To report adverse events or product complaints
Technical Support	1-800-241-4002 (M-F, 8:00 am – 6:00 pm CT)	Additional information or technical assistance

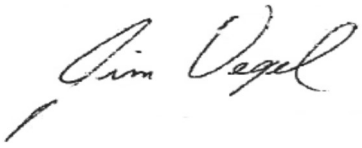
The U.S. Food and Drug Administration (FDA) has been notified of this action.

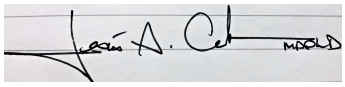
Adverse reactions or quality problems experienced with the use of this product may be reported to the FDA's MedWatch Adverse Event Reporting program either online, by regular mail, or by fax.

- Complete and submit the report **Online**: www.fda.gov/medwatch
- **Regular Mail or Fax**: Download the form at www.fda.gov/medwatch or call 1-800-332-1088 to request a reporting form, then complete and return to the address on the pre-addressed form or submit by fax to 1-800-FDA-0178

ICU Medical is committed to patient safety and is focused on providing exceptional product reliability and the highest level of customer satisfaction. Thank you for your prompt support on this important matter. We appreciate your cooperation.

Sincerely,





Jim Vegel
Vice President, Quality

Dr. Jesus Cabrera
Chief Medical Officer

Enclosures:

- *Customer Response Form*