

[Month DD, YYYY]

via [INSERT METHOD]

**URGENT FIELD SAFETY NOTICE  
MEDICAL DEVICE CORRECTION**

**Datascope Cardiosave Hybrid and Rescue Intra-Aortic Balloon Pumps (IABP)  
Cybersecurity Vulnerabilities– Ripple20**

AFFECTED PRODUCT	PART NUMBER	DISTRIBUTION DATE
Cardiosave Hybrid IABP Cardiosave Rescue IABP	All	All

**PLEASE FORWARD THIS INFORMATION TO ALL CURRENT AND POTENTIAL CARDIOSAVE HYBRID and CARDIOSAVE RESCUE IABP USERS WITHIN YOUR HOSPITAL / FACILITY.**

**IF YOU ARE A DISTRIBUTOR WHO HAS SHIPPED ANY AFFECTED PRODUCTS TO CUSTOMERS, PLEASE FORWARD THIS DOCUMENT TO THEIR ATTENTION FOR APPROPRIATE ACTION.**

Dear **Hospital Contact,**

Datascope/Getinge is initiating a voluntary Medical Device Correction for the Cardiosave Hybrid and Cardiosave Rescue Intra-Aortic Balloon Pump (IABP) due to cybersecurity vulnerabilities in a widely used low-level TCP/IP software library developed by Treck, Inc. that may result in a loss of communication to the Hospital Information System/Clinical Information System (HIS/CIS).

The inability to transmit therapy and waveform data from the Cardiosave IABP to the electronic record (HIS/CIS) does not impact the acute treatment of a patient on support.

Our records indicate that your facility has received one or more of the Cardiosave IABP units.

**Identification of the issue:**

On June 19th of 2020, the JSOF research lab published a series of cybersecurity vulnerabilities called Ripple20<sup>1</sup>. The publication consisted of nineteen (19) vulnerabilities that affects hundreds of millions of Ethernet/Internet connection capable devices.

<sup>1</sup> <https://www.jsof-tech.com/ripple20/>

Getinge’s investigation revealed that five (5) of the nineteen (19) vulnerabilities may affect the operating system in Cardiosave IABP devices. If any of the vulnerabilities were exploited, Ethernet communication would be lost, and the Cardiosave will not be able to communicate to the Hospital Information System/Clinical Information System (HIS/CIS) to send therapy and waveform data.

Although it will not send therapy and waveform data to the HIS/CIS, the Cardiosave IABP will still deliver therapy to the patient as intended, and there will be no degradation in performance.

The five vulnerabilities are listed in the table below:

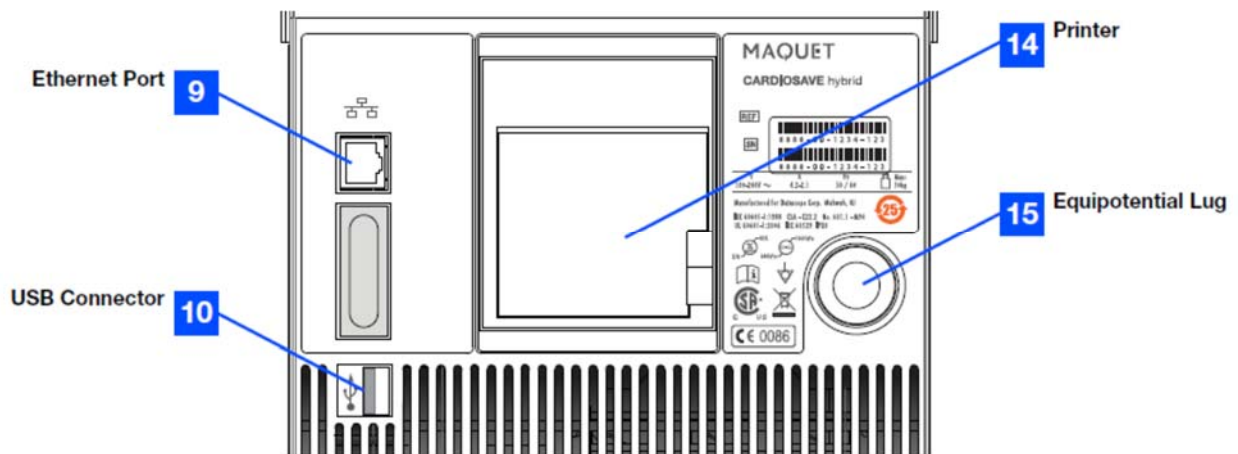
Vulnerability	Details
CVE-2020-11896	Improper handling of length parameter in IP4/UDP.
CVE-2020-11906	Improper Input Validation in Ethernet Link Layer component
CVE-2020-11907	Improper handling of length parameter inconsistency in TCP component
CVE-2020-11911	Improper access control in ICMPv4.
CVE-2020-11914	Improper input validation in ARP component

It is important to note that there have been no adverse events or deaths attributed to this issue.

**Interim Immediate actions to be taken by User:**

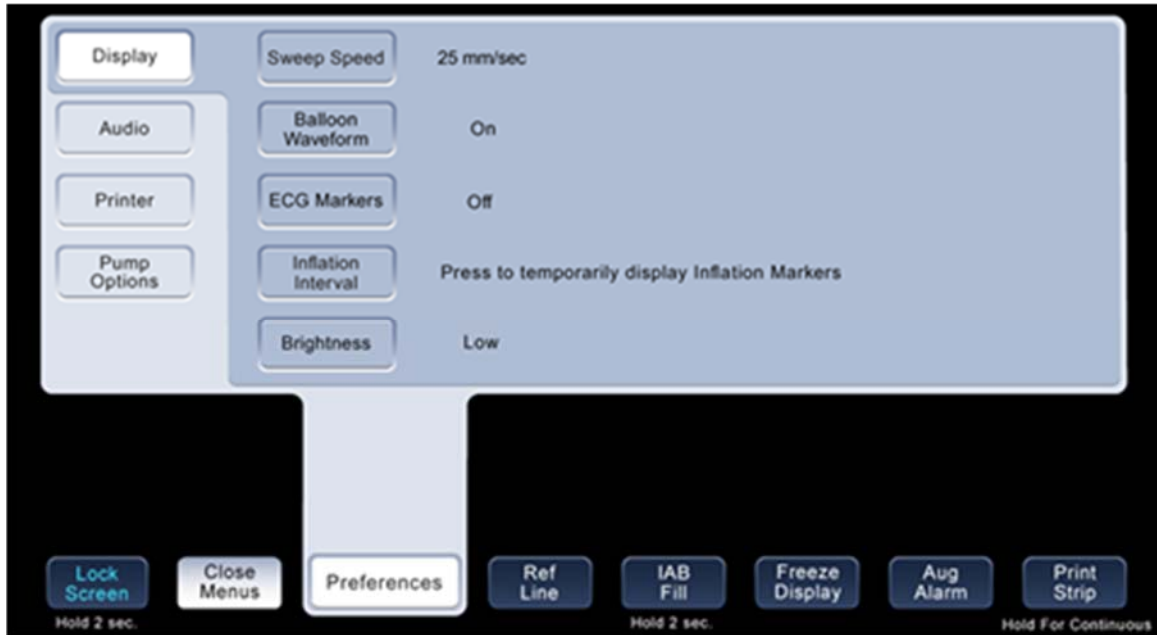
To ensure that Cardiosave Hybrid or Cardiosave Rescue are not susceptible to the Ripple20 vulnerabilities, users can disconnect the Ethernet cable from the Cardiosave Ethernet Port identified as item 9 in the image in Figure 1 below:

Figure 1 (back of Cardiosave Hybrid and Cardiosave Rescue unit)

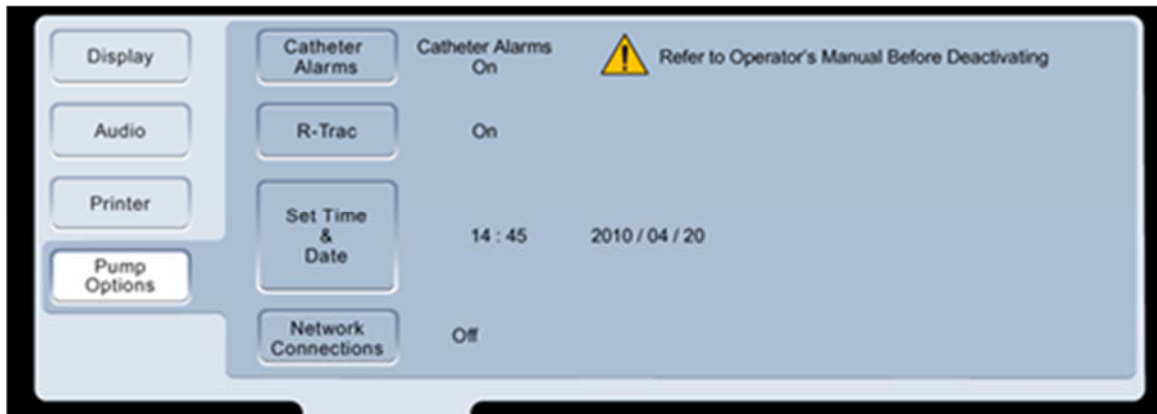


Additionally the user can turn off Network Connections via the **Network Connections** settings in the **Pump Options** menu. Ensure that the Connection Status indicator displays red after the Network Connections are set to Off.

To access the Network Settings Menus first press the **Preferences** key on the bottom row of the Keypad display to display the **Preferences Menu**.



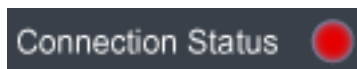
Within the **Preferences Menu** select the **Pump Options** key to open the **Pump Options** sub-menu.



With the **Pump Options** Menu open select the **Network Connections** key to access the network options.



Select the **Off** key and confirm that the **Connection Status** indicator displays red.



These actions will isolate Cardiosave from any potential external network based vulnerabilities. Cardiosave does not support any other network connection types other than the direct connected Ethernet cables.

**Corrective Action:**

Datascope/Getinge is currently developing a software correction to address this issue. A Datascope/Getinge service representative will contact you to schedule the installation of the updated software. This work will be done at no cost to your facility.

Please complete and sign the attached URGENT FIELD SAFETY NOTICE MEDICAL DEVICE CORRECTION– RESPONSE FORM (page 5) to acknowledge that you have received this notification. Return the completed form to Datascope/Getinge by e-mailing a scanned copy to **INSERT LOCAL SSU EMAIL ADDRESS** or by faxing the form to **INSERT LOCAL SSU FAX NUMBER**.

We apologize for any inconvenience this Medical Device Correction may cause. If you have any questions, please contact your local Datascope/Getinge representative.

This notification is being made with the knowledge of the U.S. Food and Drug Administration.

Sincerely,

**Recall Coordinator Name**  
**Recall Coordinator Title, Regulatory Affairs and Field Action Compliance**  
Getinge

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**URGENT FIELD SAFETY NOTICE  
MEDICAL DEVICE CORRECTION  
RESPONSE FORM**

**Datascope Cardiosave Hybrid and Rescue Intra-Aortic Balloon Pumps (IABP)  
Cybersecurity Vulnerabilities– Ripple20**

**FAX BACK TO:** INSERT LOCAL SSU FAX NUMBER **or EMAIL TO:** INSERT LOCAL SSU EMAIL ADDRESS

**ADD ACCOUNT#  
[FACILITY NAME  
STREET ADDRESS  
CITY, STATE, ZIP CODE]**

I acknowledge that I have reviewed and understand this Urgent Medical Device Correction Letter for the affected Cardiosave Intra-Aortic Balloon Pump(s) at this facility.

I confirm that all users of the Cardiosave Intra-Aortic Balloon Pump(s) at this facility have been notified accordingly.

Please provide the required information and signature below.

Facility Representative:

Signature: \_\_\_\_\_ Date: \_\_\_\_\_

Name: \_\_\_\_\_ Phone: \_\_\_\_\_

Title: \_\_\_\_\_ Department: \_\_\_\_\_

Hospital Name: \_\_\_\_\_

Address, City and State: \_\_\_\_\_

**Return the completed form by FAX to** INSERT LOCAL SSU FAX NUMBER **or by EMAIL to** INSERT LOCAL SSU EMAIL ADDRESS