

Welch Allyn, Inc. 4341 State Street Road Skaneateles Falls, NY 13153 USA	URGENT: Field Safety Notice	MOD1319
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Date

WA-MOD1319-XXXX
Customer name
Customer Street Address
Customer Country, Zip

Commercial name of affected product:
Welch Allyn ELI380 RESTING ELECTROCARDIOGRAPH

Type of action: Field Safety Corrective Action – Software Upgrade

Dear Welch Allyn Customer,

Description of the problem:

The ELI® 380 is a 12-Lead/15-Lead Resting Electrocardiograph indicated for use to acquire, analyze, display, and print electrocardiograms. Under certain circumstances, the ELI380 may lose wireless connection to the health care facility network. An Internal investigation confirmed that the radio within the device can become disassociated with the wireless access point after the ECG is idle for 30 or more minutes in an environment with high network congestion. The issue is caused by a software fault, isolated to ELI380 products utilizing the Laird WLAN module and operating with software version 2.4.1 or 2.4.2.

Potential Risk:

If the ELI380 Resting Electrocardiograph remains idle for greater than 30 minutes in an environment with high network congestion, the device may lose wireless connection to the network. An error message will be displayed to the user, indicating the device has lost the network connection. The device has a STAT ECG button that is intended to acquire an emergency ECG, and a wireless connection is not needed to perform this function. If an ECG test is not performed using the STAT ECG function, there could be a potential delay in critical care/diagnosis. Welch Allyn's internal risk assessment indicates that any level of patient harm is improbable. To date, there have been more than 1,344,034 estimated patient experiences with the affected units and Welch Allyn has received one complaint of a delay in critical patient care.

Affected Product:

The products associated with this Field Safety Notice were manufactured between June 03, 2019 and July 27, 2020. A list of the affected part numbers is provided in Table 1.

Action to be taken by the user:

- If an ECG test is needed when there is no wireless connection, follow the user manual and acquire an ECG using the STAT ECG mode.

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- The operating software installed on the ELI380 Resting Electrocardiograph requires to be upgraded to version 2.4.3 in order to address this issue. Included with this Field Safety Notice is a USB flash drive with the update software to be used to update your affected units identified in Appendix C.
- Enclosed in Appendix C is a list of impacted devices and serial numbers. Locate the devices in your possession and upgrade the software as soon as possible following the instructions in Appendix B or using the instructions provided on the USB flash drive. After the software is successfully upgraded, complete the enclosed Field Safety Notice Confirmation Form in Appendix C. Sign and return the form to mod1319@hillrom.com.

Note: Instructions are provided in Appendix A to locate the serial number on the device if needed.

- If technical assistance is needed during the installation process, please contact the appropriate regional technical support via phone or email using the contact information provided in the "Contact Reference Person" table below.
- This notice should be sent to all members of your organization who need to be aware and/or to any organization where the potentially affected devices have been transferred.

Contact Reference Person:

Should you have any questions regarding this notification, please contact Hillrom/Welch Allyn Technical Support, using email or number below.

Region/Country	Technical Support Phone	Technical Support Email
CZECH REPUBLIC	+353 (0) 46 90 67 790, Option 3	eme.techsupport@hillrom.com
FINLAND	+358 969 379 386	eme.techsupport@hillrom.com
GERMANY	+49 6950 985 132, Option 3	eme.techsupport@hillrom.com
HUNGARY	+353 (0) 46 90 67 790, Option 3	eme.techsupport@hillrom.com
ITALY	+39 0269682425, Option 3	eme.techsupport@hillrom.com
ISRAEL	+353 (0) 46 90 67 790, Option 3	eme.techsupport@hillrom.com
MIDDLE EAST	+353 46 90 67790, Option 3	eme.techsupport@hillrom.com
NETHERLANDS	+31 (0) 20 206 13 60, Option 3	eme.techsupport@hillrom.com
NORWAY	+47 23 16 25 27	eme.techsupport@hillrom.com
SWITZERLAND	+41 44 6545315	eme.techsupport@hillrom.com
UK	+44 207 365 6780, Option 3	eme.techsupport@hillrom.com

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Please ensure this notice is circulated to all appropriate personnel. This may include, but is not limited to:

• A&E departments	• In-house maintenance staff
• Adult intensive care units	• IV nurse specialists
• All wards & Clinics	• Medical directors
• Biomedical engineering staff	• Nursing executive directors
• Clinical governance leads	• Oncology units
• Day case theatres	• Pediatric intensive care units
• EBME departments	• Risk managers
• Equipment stores & Libraries	• Supplies managers
• Health and safety managers	• Theatres

The undersign confirms that this notice has been communicated to your local Regulatory Agency.

Sincerely,



Mark Elliott
 Director, Quality Assurance

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Table 1: Affected Product

Part Number	Description	Device Identifier
ELI380-ABS12	ELI380 LCD WAM WLAN USB SEC IEC BAN	00732094265545
ELI380-ABX12	ELI380 LCD WAM WLAN USB IEC BAN	00732094265514
ELI380-ACQ12	ELI380 LCD WAM WLAN USBDICOM PDQ SEC IEC	00732094319439
ELI380-ACS11	ELI380 LCD WAM WLAN USB DICOM SEC AHA B	00812345026313
ELI380-ACX11	ELI380 LCD WAM WLAN USB DICOM AHA BAN	00812345024531
ELI380-ACX12	ELI380 LCD WAM WLAN USB DICOM IEC BAN	00732094265439
ELI380-ACX21	ELI380 LCD AM12 WLAN USB DICOM AHA BAN	00812345024562
ELI380-ACX22	ELI380 LCD AM12 WLAN USB DICOM IEC BAN	00732094265422
ELI380-ACX32	ELI380 LCD AM12M WLAN USB DICOM IEC BAN	00732094265415
ELI380-DAS44	ELI380 ERGO AM15E LAN USB SEC IEC BAN	00732094265354
ELI380-DBX12	ELI380 ERGO WAM WLAN USB IEC BAN	00732094265286
ELI380-DBX22	ELI380 ERGO AM12 WLAN USB IEC BAN	00732094265279
ELI380-DCP12	ELI380 ERGO WAM WLAN USB DICOM PDQ IEC B	00732094301670
ELI380-DCQ11	ELI380 ERGO WAM WLAN USB DICM PDQSEC AHA	00732094307870
ELI380-DCS11	ELI380 ERGO WAM WLAN USB DICOM SEC AHA B	00812345026917
ELI380-DCS12	ELI380 ERGO WAM WLAN USB DICOM SEC IEC B	00732094265248
ELI380-DCS21	ELI380 ERGO AM12 WLAN USB DICOM SEC AHA	00812345026924
ELI380-DCS43	ELI380 ERGO AM15E WLAN USB DICOM SEC AHA	00812345026948
ELI380-DCT12	ELI380 ERGO WAM WLAN USB DICOM SAECG IEC	00732094265217
ELI380-DCX11	ELI380 ERGO WAM WLAN USB DICOM AHA BAN	00812345026979
ELI380-DCX12	ELI380 ERGO WAM WLAN USB DICOM IEC BAN	00732094265200
ELI380-DCX21	ELI380 ERGO AM12 WLAN USB DICOM AHA BAN	00812345026986
ELI380-DCX22	ELI380 ERGO AM12 WLAN USB DICOM IEC BAN	00732094265194
ELI380-DCX32	ELI380 ERGO AM12M WLAN USB DICOM IEC BAN	00732094265187
ELI380-DCX43	ELI380 ERGO AM15E WLAN USB DICOM AHA BAN	00812345027006

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Appendix A MOD1319 Device Labeling

1. The serial number can be located on the nameplate label on the bottom side of the device and is denoted as "SN" as shown below:

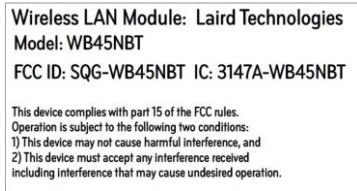


***Picture is for reference only**

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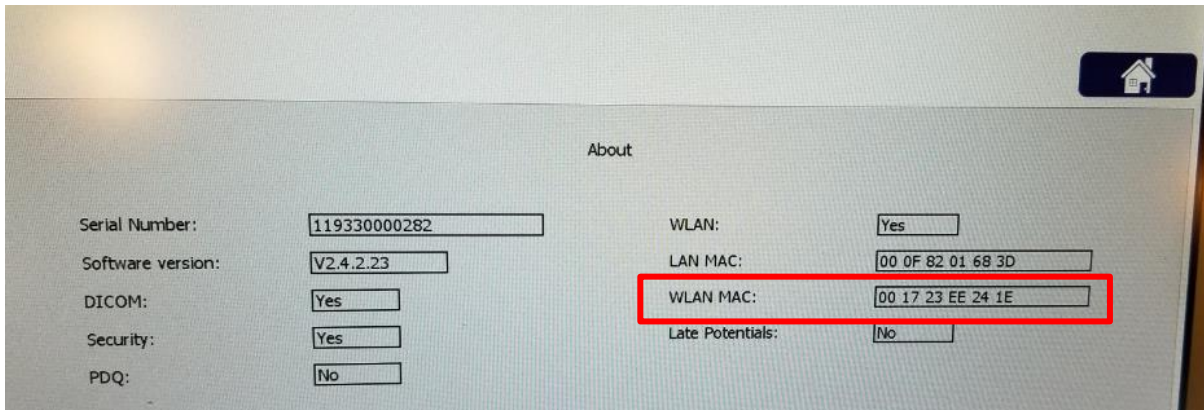
Appendix B
MOD1319 Software Installation Instructions

Version 2.4.3 can ONLY be installed on ELI380 units with a Laird WLAN module installed, as indicated by one of the following two labels being present on the bottom of the unit.



If the electrocardiograph is mounted to a transport cart and the bottom of the device is not visible, the WLAN MAC address can be used to determine the type of WLAN module installed in the unit.

The WLAN MAC address is located under the SETTINGS function of the electrocardiograph and will appear as shown below.



The WLAN MAC address contains 12 characters, where the first 6 characters are used to determine the model of the WLAN module installed. The remaining 6 digits will change with each device and is not relevant. Only upgrade units to v2.4.3 that have the MAC address shown below.

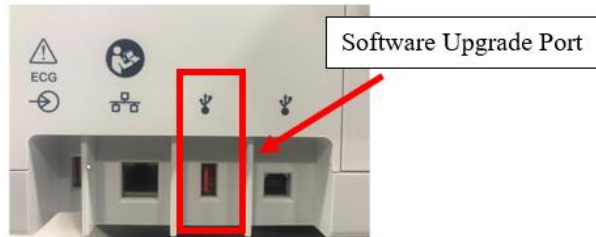
MAC address = 00 17 23 xx xx xx


As noted above, software v2.4.3 is not compatible for B&B modules. **If the MAC address is 00 0B 28 xx xx xx, do not perform the software upgrade.**

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Software Version Upgrade Process:

1. Press the On/Off (🔌) button to turn the unit off (if not already in this state).
2. Insert the USB flash drive (labeled as item # 420282) into the USB port of the unit as shown below (do not use the ECG Input port).



3. Press the On/Off button to turn the unit on.
4. Print a configuration page. Press the System Configuration button , then press Print.
5. Press **Advanced**, enter "admin" as the password (or the facility specific password).
6. Press **Service**.
7. Press **Firmware**. The display will say Reading ELI380.bin, then Erasing flash, then Writing flash, then the unit will turn off.
(if the unit does not start back up after 30 seconds, press the on/off button)

8. Press System Configuration .

9. Press **Advanced** and enter the password.

10. Press **Service**.

11. Press **Config Files**.

12. Press **Done** when the button appears, the device will then turn off.

13. Press the On/Off button to turn the unit on.

14. Press the System Configuration button .

15. Press **Print** and compare the original configuration page to the new configuration page. The new configuration page should indicate the software version v2.4.3.

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Appendix C
MOD1319 Field Safety Notice Confirmation Form

According to our records, you have received one or more impacted ELI 380 devices:

WA-MOD1319-XXXX
 Customer name
 Customer Street Address
 Customer Country, Zip

Complete, sign and return this form to mod1319@hillrom.com

1. We have reviewed and understand the attached Field Safety Notice. Yes No
2. We have reviewed the impacted serial numbers and verified all affected inventory still in our possession. Yes No
3. Check the Yes or No check box below for the serial numbers that have been upgraded to version 2.4.3.

Part Number	Device Identifier	Serial Number	Software Upgrade completed
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No

By signing below, you are confirming you have upgraded all of the impacted devices in your possession to software version 2.4.3.

Signature: _____ Date: _____

Phone Number: _____ Email Address: _____

Name of Signer (Printed) _____

Title of Signer: _____ Customer Account #: _____

Dear Distributor,

This letter is to inform you that under certain circumstances, the ELI® 380 12-Lead/15-Lead Resting Electrocardiograph may lose wireless connection to the health care facility network. The attached notification applies to all the ELI® ELI380 devices manufactured between June 03, 2019, and July 27, 2020. Our records indicate that you have purchased the affected ELI® ELI380 devices for distribution. Please distribute the attached letter to your consignees requesting they follow the instructions outlined in the communication.

If you have any questions concerning this Urgent Field Safety Notice please contact Mark Elliott, QA Director at mod1319@hillrom.com or your local Hillrom representative.

Thank you in advance for your attention on this matter.

Sincerely,



Mark Elliott
Director, Quality Assurance

Urgent Field Safety Notice - Distributor

Subject: Welch Allyn ELI380 Resting Electrocardiograph with a Laird WLAN module installed

FSCA-identifier: MOD1319

Type of action: Field Safety Corrective Action – Software Upgrade

Date: [Insert Date Here]

To: Authorized Hillrom Distributor

Details on Affected Device:

Affected devices: ELI® 380 Resting Electrocardiograph with a Laird WLAN module installed

Manufacturing Date: June 03, 2019 to July 27, 2020

A list of the affected part numbers is provided in Table 1

Background:

Description of the problem:

The ELI® 380 is a 12-Lead/15-Lead Resting Electrocardiograph indicated for use to acquire, analyze, display, and print electrocardiograms. Under certain circumstances, the ELI380 may lose wireless connection to the health care facility network. An Internal investigation confirmed that the radio within the device can become disassociated with the wireless access point after the ECG is idle for 30 or more minutes in an environment with high network congestion. The issue is caused by a software fault, isolated to ELI380 products utilizing the Laird WLAN module and operating with software version 2.4.1 or 2.4.2.

Potential Risk:

If the ELI® 380 Resting Electrocardiograph remains idle for greater than 30 minutes in an environment with high network congestion, the device may lose wireless connection to the network. An error message will be displayed to the user, indicating the device has lost the network connection. The device has a STAT ECG button that is intended to acquire an emergency ECG, and a wireless connection is not needed to perform this function. If an ECG test is not performed using the STAT ECG function, there could be a potential delay in critical care/diagnosis. Welch Allyn's internal risk assessment indicates that any level of patient harm is improbable. To date, there have been more than 1,344,034 estimated patient experiences with the affected units and Welch Allyn has received one complaint of a delay in critical patient care.

Action to be taken by Distributor:

- Enclosed in Attachment 1 is a list of impacted devices and serial numbers.
- Identify the impacted devices you have in your possession and the impacted devices you have further distributed.
- For devices still in your possession, upgrade the software to version 2.4.3 following the instructions in Appendix B of Customer Safety Notice Section or using the instructions provided on the USB flash drive.



- For devices you have further distributed, mail the enclosed Customer Field Safety Notice, with a list of impacted devices for each customer (you will need to pull the specific products shipped out of Attachment 1 and move them to Appendix C (Customer Safety Notice Section) specific to what you sent to each of your accounts. Appendix C in the Customer Safety Notice Section will be different for each account).
- The Customer Field Safety Notice documents will be: Customer Letter, Table 1, Appendix A (Device Labelling), Appendix B (Software Installation Instructions), Appendix C (Confirmation Form specific by account), and Appendix D (with your contact information added).
- Collect the signed Field Safety Notice Confirmation Forms from your customers.
- After the software is successfully upgraded for all the impacted devices in your possession and/or in your consignee's possession, complete the enclosed **MOD1319 Distributor Field Safety Notice Confirmation Form** in Attachment 1. Sign and return the form to mod1319@hillrom.com

Contact reference person:

If you have any questions regarding this safety notice, please contact Mark Elliott, QA Director at mod1319@hillrom.com or Hill-Rom Technical Support, or your Hill-Rom representative.

Yours sincerely

Hill-Rom Technical Support

List of Attachments:

Attachment 1 - MOD1319 Distributor Field Safety Notice Confirmation Form
Attachment 2 - Customer Field Safety Notice Section

List of Appendices:

Appendix A - MOD1319 Device Labelling
Appendix B - MOD1319 Software Installation Instructions
Appendix C - MOD1319 Customer Medical Device Correction Response Form
Appendix D - Authorized Hillrom Medical Distributor Information

Table 1: Affected Product

Part Number	Description	Device Identifier
ELI380-ABS12	ELI380 LCD WAM WLAN USB SEC IEC BAN	00732094265545
ELI380-ABX12	ELI380 LCD WAM WLAN USB IEC BAN	00732094265514
ELI380-ACQ12	ELI380 LCD WAM WLAN USBDICOM PDQ SEC IEC	00732094319439
ELI380-ACS11	ELI380 LCD WAM WLAN USB DICOM SEC AHA B	00812345026313
ELI380-ACX11	ELI380 LCD WAM WLAN USB DICOM AHA BAN	00812345024531
ELI380-ACX12	ELI380 LCD WAM WLAN USB DICOM IEC BAN	00732094265439
ELI380-ACX21	ELI380 LCD AM12 WLAN USB DICOM AHA BAN	00812345024562
ELI380-ACX22	ELI380 LCD AM12 WLAN USB DICOM IEC BAN	00732094265422
ELI380-ACX32	ELI380 LCD AM12M WLAN USB DICOM IEC BAN	00732094265415
ELI380-DAS44	ELI380 ERGO AM15E LAN USB SEC IEC BAN	00732094265354
ELI380-DBX12	ELI380 ERGO WAM WLAN USB IEC BAN	00732094265286
ELI380-DBX22	ELI380 ERGO AM12 WLAN USB IEC BAN	00732094265279
ELI380-DCP12	ELI380 ERGO WAM WLAN USB DICOM PDQ IEC B	00732094301670
ELI380-DCQ11	ELI380 ERGO WAM WLAN USB DICM PDQSEC AHA	00732094307870
ELI380-DCS11	ELI380 ERGO WAM WLAN USB DICOM SEC AHA B	00812345026917
ELI380-DCS12	ELI380 ERGO WAM WLAN USB DICOM SEC IEC B	00732094265248
ELI380-DCS21	ELI380 ERGO AM12 WLAN USB DICOM SEC AHA	00812345026924
ELI380-DCS43	ELI380 ERGO AM15E WLAN USB DICOM SEC AHA	00812345026948
ELI380-DCT12	ELI380 ERGO WAM WLAN USB DICOM SAECG IEC	00732094265217
ELI380-DCX11	ELI380 ERGO WAM WLAN USB DICOM AHA BAN	00812345026979
ELI380-DCX12	ELI380 ERGO WAM WLAN USB DICOM IEC BAN	00732094265200
ELI380-DCX21	ELI380 ERGO AM12 WLAN USB DICOM AHA BAN	00812345026986
ELI380-DCX22	ELI380 ERGO AM12 WLAN USB DICOM IEC BAN	00732094265194
ELI380-DCX32	ELI380 ERGO AM12M WLAN USB DICOM IEC BAN	00732094265187
ELI380-DCX43	ELI380 ERGO AM15E WLAN USB DICOM AHA BAN	00812345027006



**Attachment 1 - Distributor
MOD1319 Distributor Field Safety Notice Confirmation Form**

According to our records, you have received one or more impacted ELI 380 devices.

Complete, sign and return this form to mod1319@hillrom.com

1. We have reviewed and understand the attached Field Safety Notice. Yes No
2. We have reviewed the impacted serial numbers and verified all affected inventory still in our possession and the units sold to our consignees. Yes No
3. Check the Yes or No check box below for the serial numbers that have been upgraded to version 2.4.3.

Part Number	Device Identifier	Serial Number	Software Upgrade completed
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No

By signing below, you are confirming you have upgraded all of the impacted devices in your possession and/or in your consignee's possession to software version 2.4.3.

Signature: _____ Date: _____

Phone Number: _____ Email Address: _____

Name of Signer (Printed) _____

Title of Signer: _____

Distributor Company Name: _____

Attachment 2

Customer Field Safety Notice Section

Urgent Field Safety Notice - Customer

Subject: Welch Allyn ELI380 Resting Electrocardiograph with a Laird WLAN module installed

FSCA-identifier: MOD1319

Type of action: Field Safety Corrective Action – Software Upgrade

Date: [Insert Date Here]

To: Chief Executive, Facility Risk Manager, Facility Administrator, Facility Engineer, Vigilance Manager, Biomedical Engineering, Medical Device Liaison Officer.

Details on Affected Device:

Affected devices: ELI® 380 Resting Electrocardiograph with a Laird WLAN module installed

Manufacturing Date: June 03, 2019 to July 27, 2020

A list of the affected part numbers is provided in Table 1.

Background:

Description of the problem:

The ELI® 380 is a 12-Lead/15-Lead Resting Electrocardiograph indicated for use to acquire, analyze, display, and print electrocardiograms. Under certain circumstances, the ELI380 may lose wireless connection to the health care facility network. An Internal investigation confirmed that the radio within the device can become disassociated with the wireless access point after the ECG is idle for 30 or more minutes in an environment with high network congestion. The issue is caused by a software fault, isolated to ELI380 products utilizing the Laird WLAN module and operating with software version 2.4.1 or 2.4.2.

Potential Risk:

If the ELI380 Resting Electrocardiograph remains idle for greater than 30 minutes in an environment with high network congestion, the device may lose wireless connection to the network. An error message will be displayed to the user, indicating the device has lost the network connection. The device has a STAT ECG button that is intended to acquire an emergency ECG, and a wireless connection is not needed to perform this function. If an ECG test is not performed using the STAT ECG function, there could be a potential delay in critical care/diagnosis. Welch Allyn's internal risk assessment indicates that any level of patient harm is improbable. To date, there have been more than 1,344,034 estimated patient experiences with the affected units and Welch Allyn has received one complaint of a delay in critical patient care.

Action to be taken by the user:

- If an ECG test is need when there is no wireless connection, follow the user manual and acquire an ECG using the STAT ECG mode.
- The operating software installed on the ELI380 Resting Electrocardiograph requires to be upgraded to version 2.4.3 in order to address the issue. Included with this Urgent Field Safety

DIR 80027254 Ver A

7865 North 86th Street, Milwaukee, WI 53224

Telephone 800.231.7437

www.welchallyn.com



Notice is a USB flash drive with update software to be used to update your affected units identified in Appendix C.

- o Enclosed in Appendix C is a list of impacted devices and serial numbers. Locate the devices in your possession and upgrade the software as soon as possible following the instructions in Appendix B or using the instructions provided on the USB flash drive. After the software is successfully upgraded, complete the enclosed Field Safety Notice Confirmation Form in Appendix C. Sign and return the form to your Authorized Hillrom Distributor.

Note: Instructions are provided in Appendix A to locate the serial number on the device if needed.

- o If technical assistance is needed during the installation process, please contact your Authorized Hillrom Distributor.
- o This notice should be sent to all members of your organization who need to be aware and/or to any organization where the potentially affected devices have been transferred.

Transmission of this Field Safety Notice: (if appropriate)

This notice should be sent to anyone who needs to be aware within your organization, or to any organization where the potentially affected devices have been transferred.

Hillrom confirms that the relevant Competent Authorities have been informed of this Field Safety Corrective Action.

Contact reference person:

If you have any questions regarding this safety notice, please contact your Authorized Hillrom Distributor using the contact information provided in Appendix D.

Sincerely,

Your Authorized Hillrom Distributor

List of Appendices:

Appendix A - MOD1319 Device Labelling

Appendix B - MOD1319 Software Installation Instructions

Appendix C - MOD1319 Customer Medical Device Correction Response Form

Appendix D - Authorized Hillrom Medical Distributor Information

Table 1: Affected Product

Part Number	Description	Device Identifier
ELI380-ABS12	ELI380 LCD WAM WLAN USB SEC IEC BAN	00732094265545
ELI380-ABX12	ELI380 LCD WAM WLAN USB IEC BAN	00732094265514
ELI380-ACQ12	ELI380 LCD WAM WLAN USBDICOM PDQ SEC IEC	00732094319439
ELI380-ACS11	ELI380 LCD WAM WLAN USB DICOM SEC AHA B	00812345026313
ELI380-ACX11	ELI380 LCD WAM WLAN USB DICOM AHA BAN	00812345024531
ELI380-ACX12	ELI380 LCD WAM WLAN USB DICOM IEC BAN	00732094265439
ELI380-ACX21	ELI380 LCD AM12 WLAN USB DICOM AHA BAN	00812345024562
ELI380-ACX22	ELI380 LCD AM12 WLAN USB DICOM IEC BAN	00732094265422
ELI380-ACX32	ELI380 LCD AM12M WLAN USB DICOM IEC BAN	00732094265415
ELI380-DAS44	ELI380 ERGO AM15E LAN USB SEC IEC BAN	00732094265354
ELI380-DBX12	ELI380 ERGO WAM WLAN USB IEC BAN	00732094265286
ELI380-DBX22	ELI380 ERGO AM12 WLAN USB IEC BAN	00732094265279
ELI380-DCP12	ELI380 ERGO WAM WLAN USB DICOM PDQ IEC B	00732094301670
ELI380-DCQ11	ELI380 ERGO WAM WLAN USB DICM PDQSEC AHA	00732094307870
ELI380-DCS11	ELI380 ERGO WAM WLAN USB DICOM SEC AHA B	00812345026917
ELI380-DCS12	ELI380 ERGO WAM WLAN USB DICOM SEC IEC B	00732094265248
ELI380-DCS21	ELI380 ERGO AM12 WLAN USB DICOM SEC AHA	00812345026924
ELI380-DCS43	ELI380 ERGO AM15E WLAN USB DICOM SEC AHA	00812345026948
ELI380-DCT12	ELI380 ERGO WAM WLAN USB DICOM SAECG IEC	00732094265217
ELI380-DCX11	ELI380 ERGO WAM WLAN USB DICOM AHA BAN	00812345026979
ELI380-DCX12	ELI380 ERGO WAM WLAN USB DICOM IEC BAN	00732094265200
ELI380-DCX21	ELI380 ERGO AM12 WLAN USB DICOM AHA BAN	00812345026986
ELI380-DCX22	ELI380 ERGO AM12 WLAN USB DICOM IEC BAN	00732094265194
ELI380-DCX32	ELI380 ERGO AM12M WLAN USB DICOM IEC BAN	00732094265187
ELI380-DCX43	ELI380 ERGO AM15E WLAN USB DICOM AHA BAN	00812345027006

Appendix A
MOD1319 Device Labelling

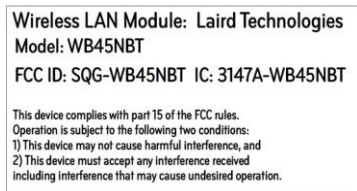
1. The serial number can be located on the nameplate label on the bottom side of the device and is denoted as "SN" as shown below:



*Picture is for reference only

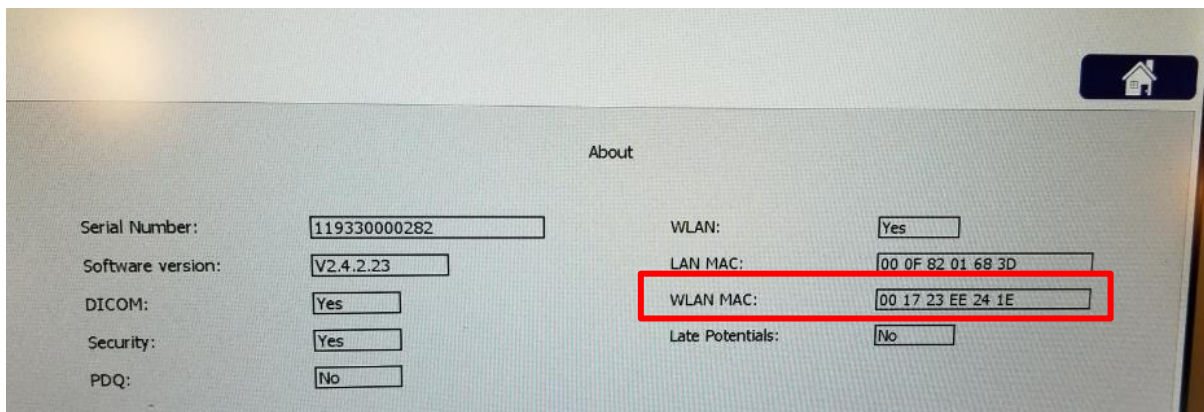
Appendix B
MOD1319 Software Installation Instructions

Version 2.4.3 can ONLY be installed on ELI380 units with a Laird WLAN module installed, as indicated by one of the following two labels being present on the bottom of the unit.



If the electrocardiograph is mounted to a transport cart and the bottom of the device is not visible, the WLAN MAC address can be used to determine the type of WLAN module installed in the unit.

The WLAN MAC address is located under the SETTINGS function of the electrocardiograph and will appear as shown below.




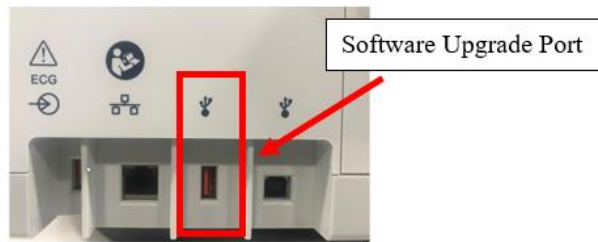
The WLAN MAC address contains 12 characters, where the first 6 characters are used to determine the model of the WLAN module installed. The remaining 6 digits will change with each device and is not relevant. Only upgrade units to v2.4.3 that have the MAC address shown below.




MAC address = 00 17 23 xx xx xx

As noted above, software v2.4.3 is not compatible for B&B modules. **If the MAC address is 00 0B 28 xx xx xx, do not perform the software upgrade.**

Software Version Upgrade Process:

1. Press the On/Off () button to turn the unit off (if not already in this state).
2. Insert the USB flash drive (labeled as item # 420282) into the USB port of the unit as shown below (do not use the ECG Input port).



3. Press the On/Off button to turn the unit on.
4. Print a configuration page. Press the System Configuration button , then press Print.
5. Press **Advanced**, enter "admin" as the password (or the facility specific password).
6. Press **Service**.
7. Press **Firmware**. The display will say Reading ELI380.bin, then Erasing flash, then Writing flash, then the unit will turn off.
(if the unit does not start back up after 30 seconds, press the on/off button)
8. Press System Configuration .
9. Press **Advanced** and enter the password.
10. Press **Service**.
11. Press **Config Files**.
12. Press **Done** when the button appears, the device will then turn off.
13. Press the On/Off button to turn the unit on.
14. Press the System Configuration button .
15. Press **Print** and compare the original configuration page to the new configuration page. The new configuration page should indicate the software version v2.4.3.



Appendix C
MOD1319 Customer Field Safety Notice Confirmation Form

According to our records, you have received one or more impacted ELI 380 devices.

Complete, sign and return this form to your Authorized Hillrom distributor.

1. We have reviewed and understand the attached Field Safety Notice. Yes No
2. We have reviewed the impacted serial numbers and verified all affected inventory still in our possession. Yes No
3. Check the Yes or No check box below for the serial numbers that have been upgraded to version 2.4.3.

Part Number	Device Identifier	Serial Number	Software Upgrade completed
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No

By signing below, you are confirming you have upgraded all of the impacted devices in your possession to software version 2.4.3.

Signature: _____ Date: _____

Phone Number: _____ Email Address: _____

Name of Signer (Printed) _____

Title of Signer: _____ Facility Name: _____



Appendix – D
Authorized Hillrom Medical Distributor Information

Distributor Company:

Contact Person:

Phone:

Email: