

Alaris[®] System Maintenance v9.5.X

Software User Manual

Model 8975, v9.5.X
Guardrails[®] Suite MX
Compatible with v9.X Alaris[®] PC units



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Alaris® System
Maintenance v9.5.X
Software User Manual
Part Number: 11614346

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Chapter 1

Introduction

Defined Terms

The following table identifies the defined terms used throughout this document for certain trademarked products and product features.

Product/Feature	Defined Term
Alaris® Auto-ID module	Auto-ID Module
Alaris® EtCO ₂ module	EtCO ₂ Module
Alaris® PCA module	PCA Module
Alaris® PC unit	PC Unit
Alaris® Pump module	Pump Module
Alaris® SpO ₂ module	SpO ₂ Module
Alaris® Syringe module	Syringe Module
Alaris® System Maintenance	System Maintenance Software
Guardrails® alert	Alert
Guardrails® clinical advisories	Clinical Advisories
Guardrails® CQI report	CQI Report
Guardrails® data set	Data Set
Guardrails® Editor software	Editor Software

About the Software

This software is provided under and subject to a license from CareFusion.

System Maintenance Software version 9.5.X allows you to perform routine maintenance on an Alaris® System. The functionality of this software replaces maintenance software (MSW) and the Flash tool for the Alaris® System. To use this software, you install it on a personal computer and then connect a serial cable between the personal computer and a PC Unit.

The System Maintenance software kit includes:

- System Maintenance software CD-ROM
- Serial cable
- This Software User Manual (pdf on CD-ROM and printed version)

A USB serial adapter (not supplied) can be used to connect one or more PC Units to a personal computer equipped with System Maintenance Software. The USB serial adapter must meet the following specifications:

- Plugs into a USB port on a personal computer.
- Provides RS-232 male DB9 port(s) for direct connection to PC Units.
- Draws its power from USB connection—a power adapter is not required.

Feature Summary

System Maintenance software can be used to perform the following functions:

- Maintain the PC Unit.
- Upgrade firmware on the PC Unit and attached modules.
- Perform preventive maintenance/check-in.
- Set and check preventive maintenance reminder dates.
- Troubleshoot and repair instruments.
- Module Calibration, as applicable.
- Set up, test, and configure the Auto-ID Module.
- Download and view instrument logs.
- Download CQI data logs.
- Transfer Data Sets from the personal computer to the PC Unit.
- Configure wireless network settings.
- Test communications board interaction with a wireless network.
- Disable Wireless Network configuration option.
- Support wireless protocols for b, b/g, a/b/g, and d networks.
- Support wireless security for WEP, WPA, and WPA2.

Perform all pre-inspections before instrument use. A qualified technician or biomedical engineer must perform inspections at least once a year or as required in accordance with CareFusion guidelines.

NOTE

- Inspections are intended to comply with requirements specified by the Joint Commission on Accreditation of Healthcare Organizations (JCAHO).
- You can purchase a service agreement from CareFusion for preventive maintenance inspections.

Warnings

Warnings provide information needed to safely and effectively use the Alaris® product.

A **WARNING** is an alert to a potential hazard which could result in serious personal injury and/or product damage if proper procedures are not followed.

WARNING

- **At no time should the System Maintenance software be used while the Alaris® System is connected to a patient.**
- **System Maintenance software and procedures are intended to be performed only by qualified personnel, or under direct supervision by qualified personnel.**
- **Prior to use, CareFusion recommends that you become familiar with each instrument and accessory used with the System Maintenance software. For specific instrument warnings, cautions, inspection requirements, and other related instrument information, refer to the applicable Alaris® System DFU and/or service manual.**
- **The instrument case should only be opened by qualified service personnel using proper grounding techniques. When the instrument case is opened, electrical shock hazard exists which can result in serious injury to persons and instrument component damage.**
- **To ensure proper calibration, run the post tests for the Pump Module and the accuracy tests for the Syringe Module after calibration. These tests are displayed in the default Calibration test lists.**
- **Ensure that the instrument is in operational status before returning it to patient use.**
- **Your PC Unit might not be compatible with every module. To determine module compatibility, refer to the applicable Alaris® System DFU.**

Cautions

Cautions provide information needed to safely and effectively use the Alaris® product.

A **CAUTION** is an alert to a potential hazard which could result in minor personal injury and/or product damage if proper procedures are not followed.

CAUTION

- Loading the System Maintenance Software is considered a non-clinical service activity. Interconnecting a medical device with a personal computer might cause the safety or electromagnetic environment to change while the connection exists. The threat of higher leakage currents or EMI disturbance levels might be present. Remove the connection at both ends before patient use.
- By default, the System Maintenance Software is set to automatically clear logs after they are downloaded. Once log entries have been cleared, they cannot be downloaded again from the instrument. The log entries remain in the database and can be displayed using the View Reports tab in the main window.
- Only one PC Unit should be connected to a group of attached modules at any given time. Connection of multiple PC Units to each other will cause damage to the IUI connectors.
- Only one SpO₂ Module can be connected to the PC Unit at any one time.
- If a Pump Module fails the patient-side occlusion or rate accuracy verification (RAV) tests, calibration is required. Failure to calibrate and retest the instrument might result in improper instrument operation.
- For tests that require water, use distilled water at room temperature between 41°F and 104°F (5°C to 40°C). If water is not within this temperature range, the reading might be inaccurate.
- If multiple instances of the System Maintenance Software are running (typically on different personal computers), it is possible to assign different IP addresses to the same PC Unit. To ensure this does not occur, assign each PC Unit a single, unique IP address.

NOTE: If the System Maintenance Software instances are connected to the same System Maintenance Software databases, this problem cannot occur.

- Calibration verifications perform strenuous tests on the Syringe Module and PCA Module. In some cases, these tests differ from the check-in/preventive maintenance tests.
- Following calibration, use calibration verifications (rather than check-in/preventive maintenance verifications) to verify correct calibration.
- Use of accessories or cables other than those specified might result in degraded electromagnetic compatibility performance or safety of the Alaris® System.
- The flow rate out of the gas source and into the EtCO₂ Module must be greater than 0.5 liter/minute to ensure accurate readings. It is recommended to place a flow meter between the gas source and the “T” in the line (breather line).

Chapter 2

Installation and Setup

System Requirements

To run the System Maintenance software, the personal computer must meet the following minimum hardware and software requirements.

CAUTION

For optimum performance of the System Maintenance Software, ensure that the personal computer is equipped with the required hardware and software, and do not use the personal computer with unspecified accessories or cables. Use of accessories or cables other than those specified might result in degraded electromagnetic compatibility performance or safety of the Alaris® System.

Hardware Requirements

The personal computer on which you install the System Maintenance Software must meet the following requirements:

- processor, Pentium Class, 1 GHz or faster
- RAM, 512 MB minimum
- HDD (hard disk drive), 5 GB minimum of free space after the operating system is installed
- CD-ROM drive
- video card, SVGA
- monitor, minimum resolution 1024 x 768; color depth set greater than 256 colors
- port: configurable RS-232, 9-pin, serial communications; or USB
- cable, serial communications (provided)

NOTES

- If the personal computer has a *serial* port, use a serial-to-serial cable.
 - If the personal computer has a *USB* port, use a serial-to-USB cable or a serial-to-serial cable with a USB converter. USB converter products are not supported by CareFusion.
- keyboard
 - mouse
 - printer and printer connection (optional)

Software Requirements

One of the following operating systems:

- Microsoft® Windows 2000 Professional with Service Pack 4
- Microsoft® Windows XP with Service Pack 2
- Microsoft® Windows 2003 Server with Service Pack 1

Other software:

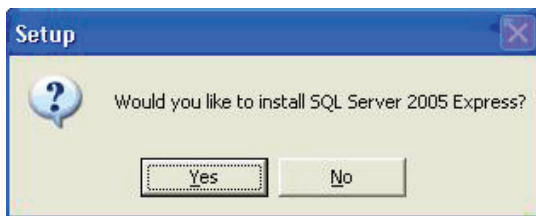
- Adobe Acrobat Reader v9.X

Install Software—Standard Installation

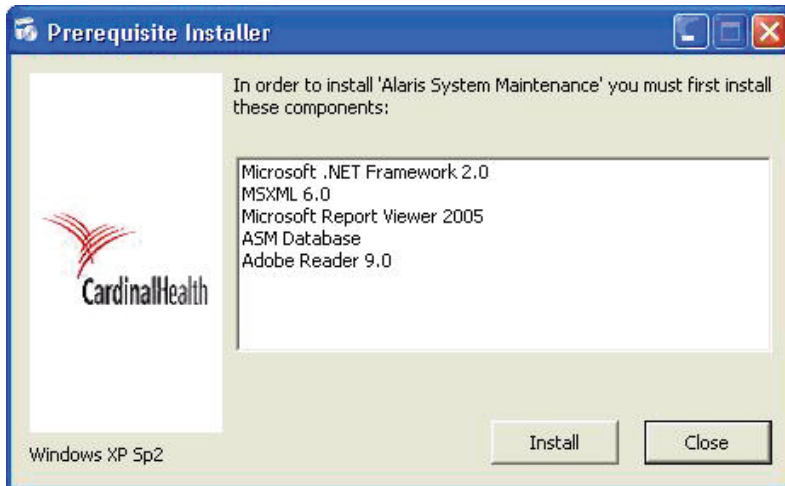
Follow these steps to install the System Maintenance Software on a personal computer for the first time. If you plan to migrate another version of the System Maintenance database, install the System Maintenance Software and migrate the data using the Migrate MSW file feature (**Options > Migrate MSW Data**).

Read all prompts carefully. During installation it is recommended that no other applications are running and that virus checking is disabled.

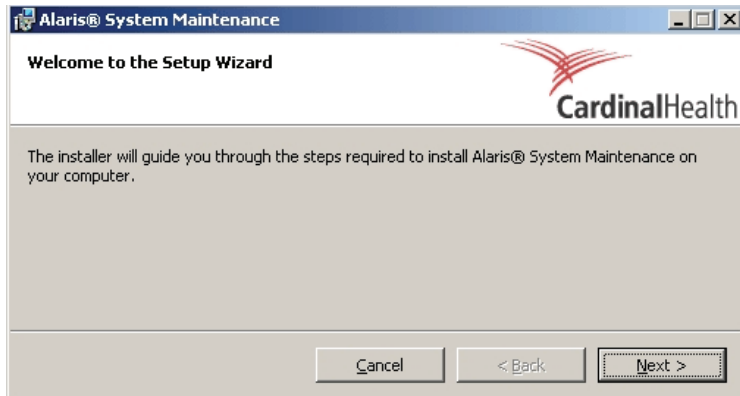
1. Insert the System Maintenance Software CD into the CD-ROM drive of the personal computer. If the installation program automatically starts, go to the next step. If the installation program does not automatically start, choose **Start > Run**, then type **d:\setup.exe** (where **d:** is CD drive). Alternatively, browse to the **Setup.exe** program on the System Maintenance Software CD, click **Open**, and then click **OK**.
2. When you are asked if you want to install SQL Server 2005 Express, click **Yes**.



3. If you are prompted to install prerequisite components, click **Install**. The components include Microsoft® .NET Framework 2.0, MSXML 6.0, Microsoft Report Viewer 2005, ASM Database (SQL Server Express 2005), and Adobe Reader 9.0.

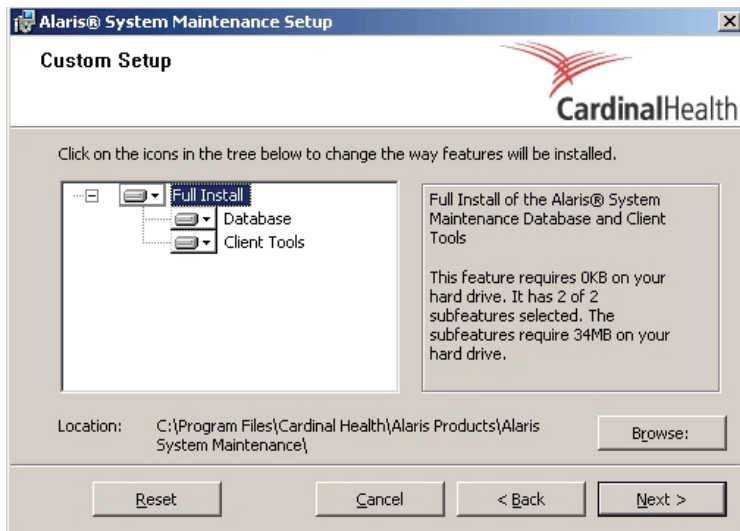


- After installing prerequisites, or if prerequisites are already installed, the installation Setup Wizard Welcome window is displayed. Click **Next**.



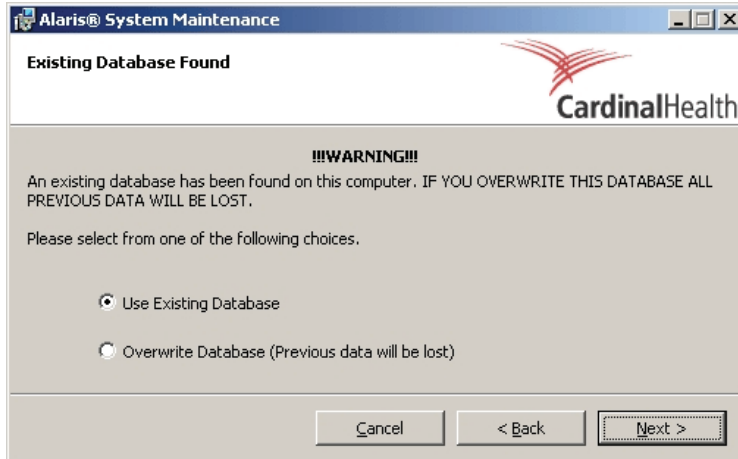
- Select the features that you want to install:
 - Full Install**—installs System Maintenance databases and client tools.
 - Database**—installs only System Maintenance databases.
 - Client Tools**—installs only System Maintenance client tools.

To install only the database, click **Client Tools** and click the red X. To install only the client tools, click **Database** and click the red X.

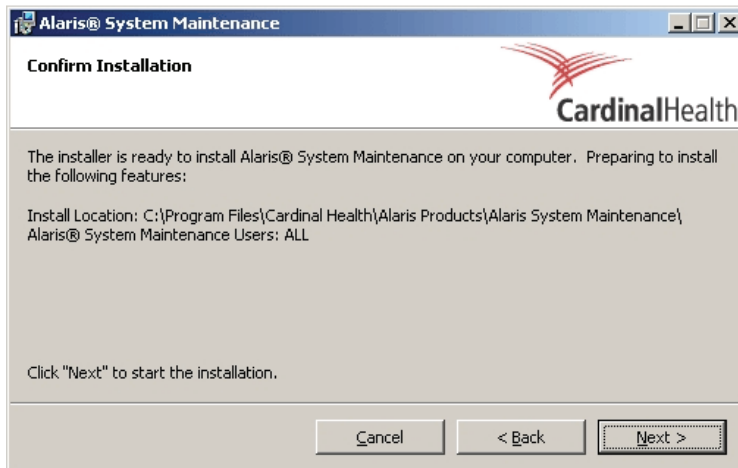


- Click **Browse** to choose a location other than the default installation directory. The default directory is **C:\Program Files\Cardinal Health\Alaris Products\Alaris System Maintenance**.

If an existing database is found, the following window is displayed. You can use the existing database (default) or overwrite the existing database (previous data will be lost). If a previous System Maintenance database was not found, this window will not be displayed.



- Click **Next** to begin installation of the components of the System Maintenance Software.
- Click **Next** to confirm installation.

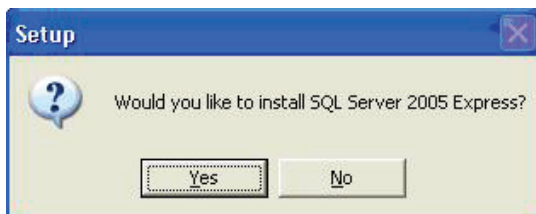


When the installation is finished, an "Installation Complete" message is displayed. It is recommended that you restart your computer if prompted to do so.

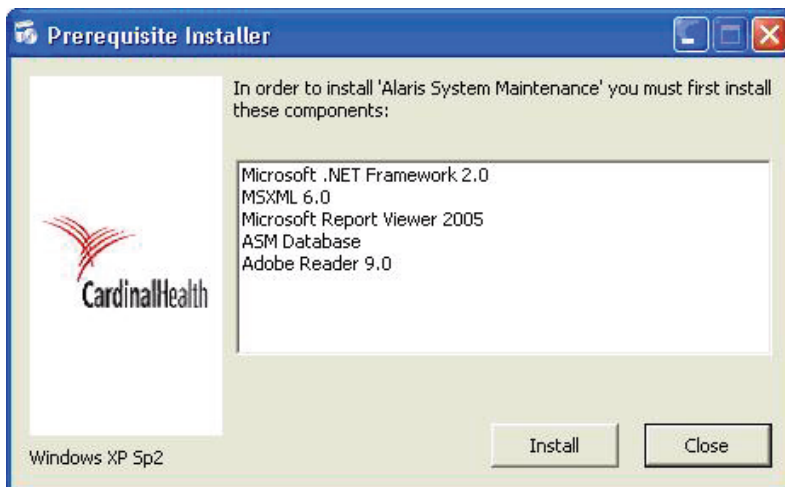
Install Software—Custom Installation

Install Centralized System Maintenance Software Database on a Dedicated Server

1. Insert the System Maintenance Software CD into the CD-ROM drive of the personal computer. If the installation program starts automatically, go to the next step. If the installation program does not start automatically, choose **Start > Run**, then type **d:\setup.exe** (where **d:** is CD drive). Alternatively, browse to the **Setup.exe** program on the System Maintenance Software CD, click **Open**, and then click **OK**.
2. When you are asked if you want to install SQL Server 2005 Express, click **Yes**.

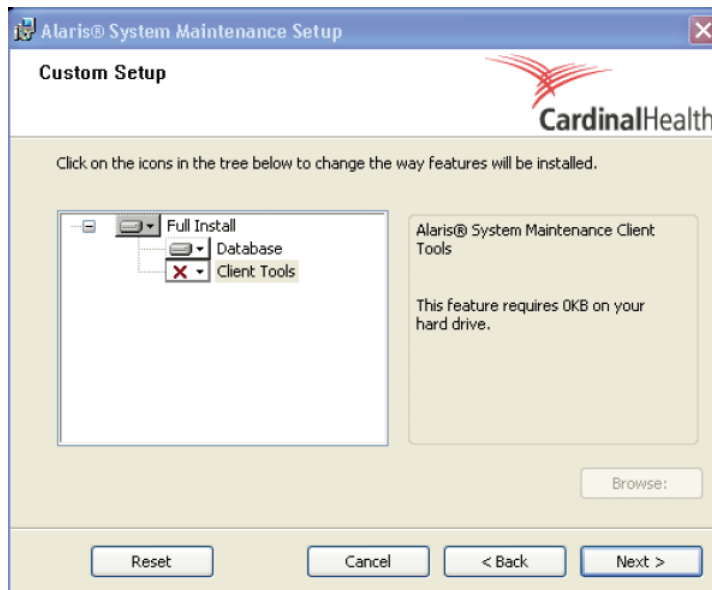


3. If you are prompted to install prerequisite components, click **Install**. The components include Microsoft .NET Framework 2.0, MSXML 6.0, Microsoft Report Viewer 2005, ASM Database (SQL Server Express 2005), and Adobe Reader 9.X.



4. Wait until the installation is completed. When the Alaris® System Maintenance welcome wizard is displayed, click **Next**.

5. On the Custom Setup window, ensure that Client Tools is *not* selected (unless you want to also install the client tools on the same computer as the System Maintenance database).

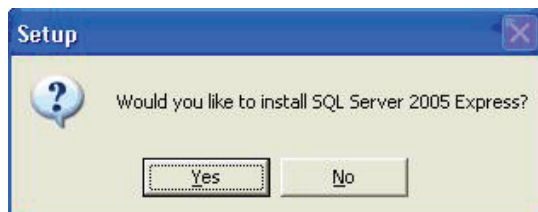


6. Click **Next**.
7. Click **Finish**.

Install Client Tools on Biomedical Technician Workstation

Follow these steps to install the client tools on the Biomedical Technician Workstation.

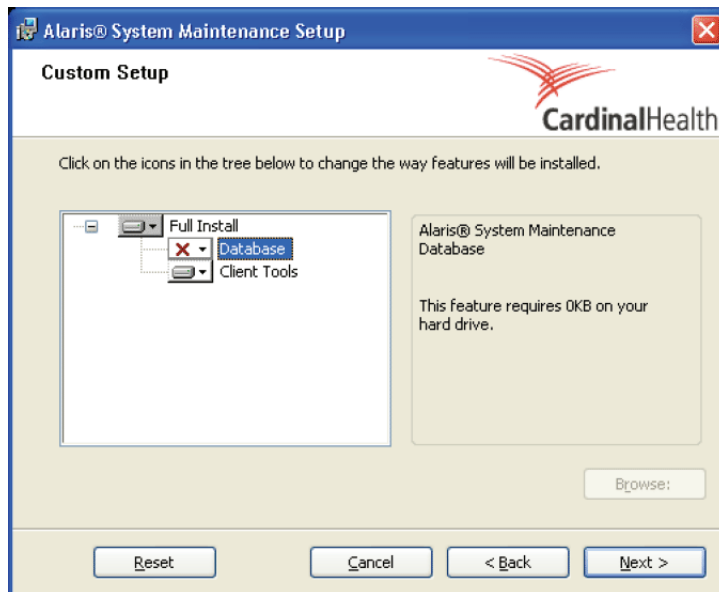
1. Insert the System Maintenance Software CD into the CD-ROM drive of the personal computer. If the installation program starts automatically, go to the next step. If the installation program does not start automatically, choose **Start > Run**, then type **d:\setup.exe** (where **d:** is CD drive). Alternatively, browse to the **Setup.exe** program on the System Maintenance Software CD, click **Open**, and then click **OK**.
2. When you are asked if you want to install SQL Server 2005 Express, click **No**.



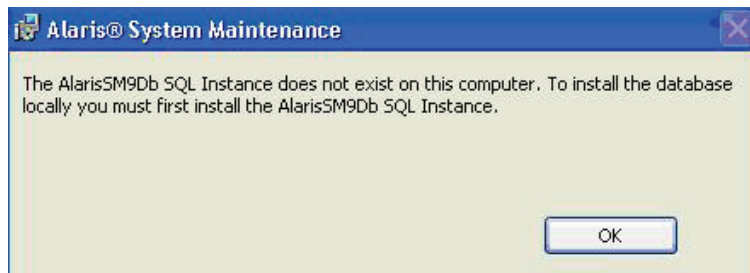
- If you are prompted to install prerequisite components, click **Install**. The components include Microsoft .NET Framework 2.0, MSXML 6.0, Microsoft Report Viewer 2005, and Adobe Reader 9.0.



- Wait until the installation is completed. When the Alaris® System Maintenance welcome wizard is displayed, click **Next**.
- On the Custom Setup window, ensure that **Database** is *not* selected.

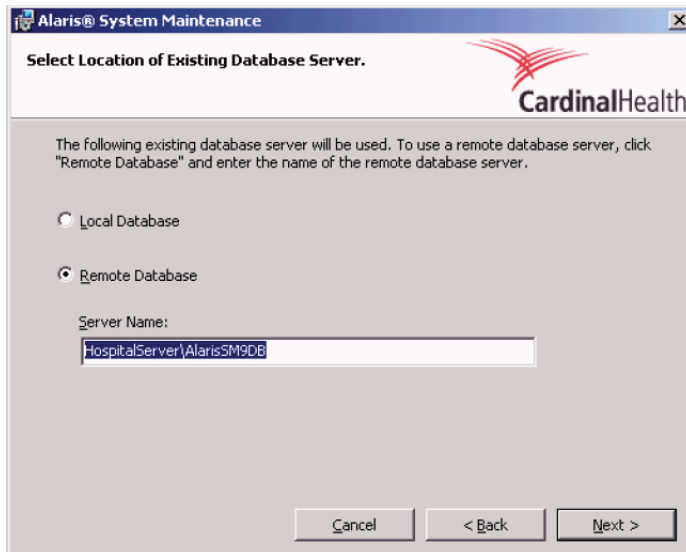


- If the Database option is selected and you chose not to install SQL Server 2005 Express, a message is displayed. Click **OK** to close the message window.



7. Click **Next**.

A window that asks you to select the database server is displayed.

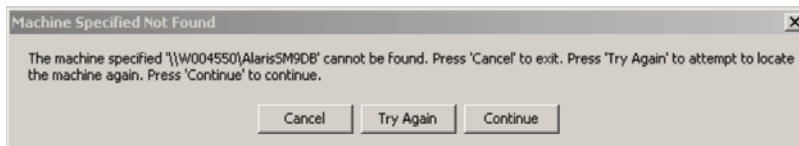


8. Choose the **Remote Database** option. Enter the server name and SQL instance.

Example: `HospitalServer\AlarissM9DB`

9. Click **Next**.

A "Machine Specified Not Found" message is displayed.



10. Click **Continue**.
11. Click **Next** to confirm.
12. Click **Finish**.

Configure Database Settings

Follow these steps to configure the database settings.

1. Start the System Maintenance Software.
2. If you are asked to select another database, click **Yes**.
3. If you are not asked, choose **Options > Database Settings** on the main menu.
The Database Settings dialog box is displayed.
4. Click **Add** to configure the connections for the Maintenance database, the Maintenance Log database, and the CQI Database.

Hospital Server

Database	Field Values
Maintenance database	SQL Server field = <HospitalServer>\AlarissM9DB Database field = Maintenance
Maintenance Log database	SQL Server field = <HospitalServer>\AlarissM9DB Database field = MaintenanceLog
CQI Database	SQL Server field = <HospitalServer>\CQI9DB Database field = CQI

Standalone Computer

Database	Field Values
Maintenance database	SQL Server field = (local)\AlarissM9Db Database field = Maintenance
Maintenance Log database	SQL Server field = (local)\AlarissM9Db Database field = MaintenanceLog
CQI Database	SQL Server field = (local)\CQI9DB Database field = CQI

NOTE

If a database does not exist on your local database server, the "(local)" SQL Service field is replaced with the SQL Server name where the database is installed.

Alaris® Server (wireless)

Database	Field Values
CQI Database	SQL Server field = <AlarisAPPServer> Database field = CQI

5. Click **Test Connection** to verify the connection to the database.

Upgrade/Reinstall Software

Before you can upgrade or reinstall the System Maintenance Software, you must uninstall the existing software.

NOTE

Uninstalling the AlarisSM9DB database is NOT required to upgrade/reinstall the software (*Uninstall AlarisSM9DB Database Instance* on page 20).

1. Uninstall the existing software (see *Uninstall Software* on page 18).
2. Install/reinstall the software (see *Install Software—Standard Installation* on page 9 or *Install Software—Custom Installation* on page 12).

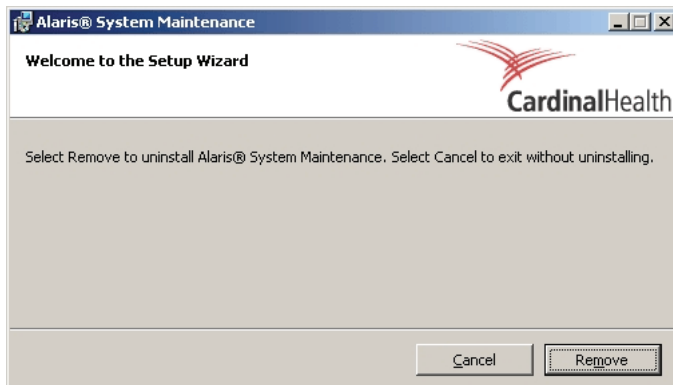
Uninstall Software

There are two ways to uninstall the System Maintenance Software. The method that you use depends on the version of software that you are uninstalling.

- System Maintenance Software CD: Use this method if the software being removed and the software that will be installed are the same software version.
- Windows Operating System: Use this method if the software being removed will be replaced with a different version of the software.

System Maintenance Software CD

1. Insert the System Maintenance Software CD into the CD-ROM drive of the personal computer. The installation program starts automatically. If the installation program does not start, choose **Start > Run**, then type **d:\Setup.exe** (where **d:** is CD drive). Alternatively, browse to the **Setup.exe** program on the System Maintenance Software CD, click **Open**, and then click **OK**.
2. Enter information about installation as prompted, including destination.
3. When the message that a previous installation has been detected is displayed, click **Remove** to completely remove the existing System Maintenance Software and its components. Existing data is preserved.



4. Clicking **Remove** displays the following confirmation window. Click **Remove** again to initiate the uninstall process.



5. When a message that indicates the software has been successfully uninstalled is displayed, click **Finish**.

Windows Operating System

1. Choose **Start > Settings > Control Panel**.
2. Double-click **Add or Remove Programs**.
3. Click **Alaris® System Maintenance**.
4. Click **Remove**.
5. Follow the prompts to complete the uninstallation procedure.

Uninstall AlarisSM9DB Database Instance

NOTE

All data will be lost when uninstalling the AlarisSM9DB database. Ensure that the database is backed up for future reference.

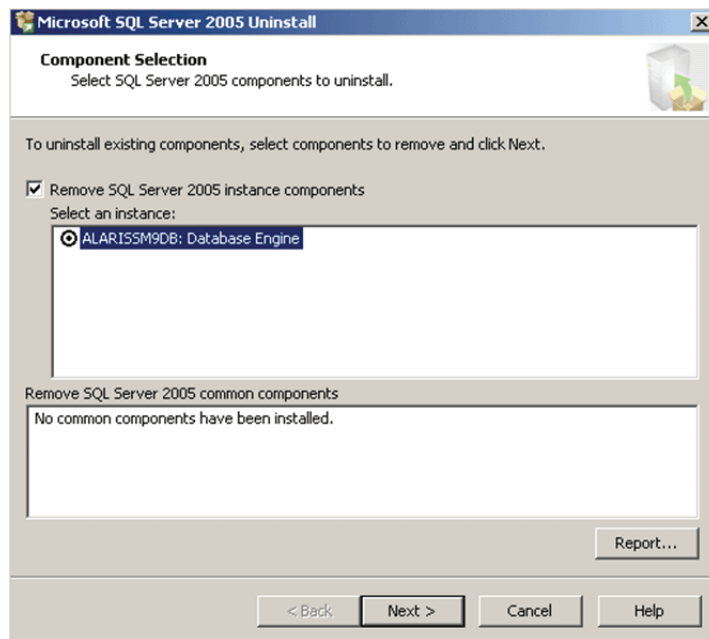
Use this procedure to uninstall the AlarisSM9DB database instance of Microsoft SQL Server 2005 Express and the following software:

- Microsoft SQL Server Native Client (if applicable)
- Microsoft SQL Server Setup Support Files (if applicable)
- Microsoft SQL Server VSS Writer (if applicable)

To uninstall the software:

1. Choose **Start > Settings > Control Panel**.
2. Double-click **Add or Remove Programs**.
3. Select the software (listed above) that you want to uninstall.
4. Click **Remove**.
5. Follow the prompts to complete the removal procedure.

If you are uninstalling Microsoft SQL Server 2005, when the following dialog box is displayed, select the instance named "AlarisSM9DB" and then click **Next** to continue with the removal procedure.



Chapter 3

Overview

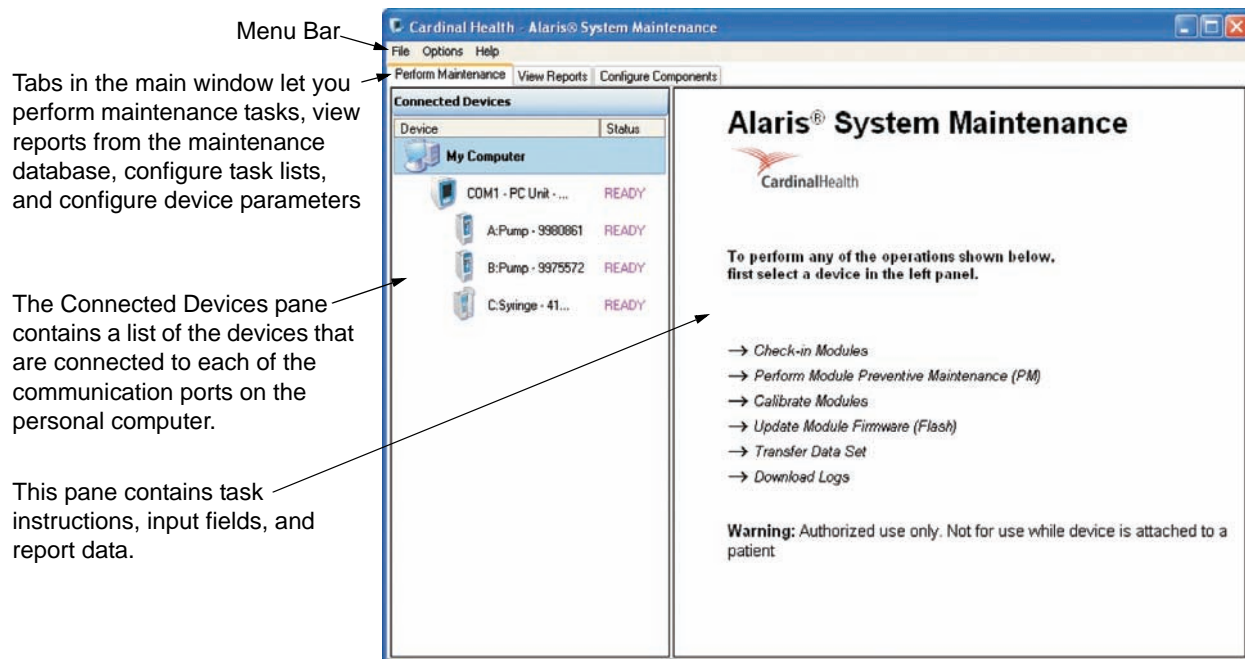
Start System Maintenance Software

1. On the personal computer desktop, double-click the Alaris® System Maintenance icon (or choose the program from the **Start** menu).

A series of status boxes is displayed as the System Maintenance Software scans for communication ports and establishes communication with connected devices.

When the startup routine is complete, the main window of the System Maintenance Software is displayed.

2. To check the version of System Maintenance Software, choose **Help > About Alaris® System Maintenance**.



Menu Bar

- **File**
 - **Refresh (F5)**—Refresh all ports.
 - **Refresh Selected Port (Ctrl + F5)**—Refresh the port that is selected in the Connected Devices pane.
 - **Exit (Alt + F4)**—Exit the software.
- **Options**
 - **Application Settings**—See *Application Settings* on page 30.
 - **Database Settings**—See *Database Settings* on page 31.
 - **Export**—Export the CQI Database as well as information for all devices that have been connected to the PC Unit.
 - **Import**—Import the CQI Database as well as information for all devices that have been connected to the PC Unit.
 - **Migrate MSW Data**—Import a previous maintenance software v8 database.
- **Help**
 - **About Alaris® System Maintenance**—View version and copyright information.

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Chapter 4

PC Unit Connection

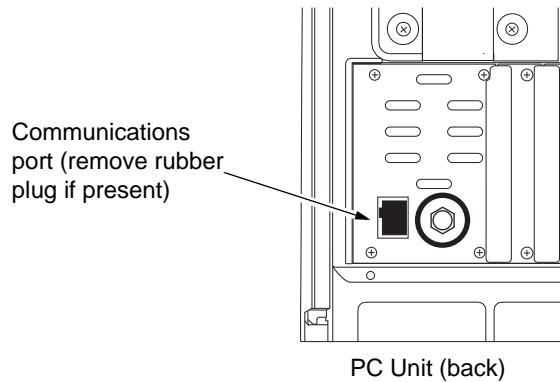
Hardware Connections

This section describes how to connect a PC Unit to the personal computer.

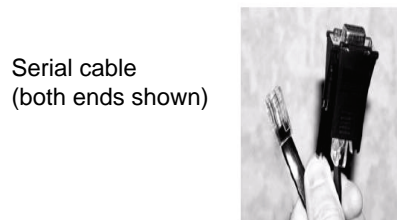
WARNING

At no time should the System Maintenance Software be used while the Alaris® System is connected to a patient.

1. If a rubber plug is covering the communications port, remove it.



2. Connect the small end of the serial cable to the communications port on the back of the PC Unit.



3. Connect the large end of the serial cable to a serial port on the personal computer.
4. If the PC Unit is not on, press the **System On** key.
5. If the System Maintenance Software is not running, start it.
6. Press F5 to refresh the ports.

Information about the PC Unit and attached modules is displayed in the Connected Devices pane.

Multiple PC Unit Connections

You can connect as many PC Units as there are available serial COM ports on the personal computer. To view all available COM ports:

1. On the main menu, click **Options > Application Settings**.
2. In the drop-down menu labeled **Show only connected ports**, click **False**.
3. Close the Application Settings window.
After a File/Refresh, a list of all available COM ports is displayed in the Connected Devices pane, whether or not they have a PC Unit attached.
4. Connect the PC Units to the available ports.
5. Press F5 to refresh the view for all ports, or press Ctrl+F5 to refresh only the port that is selected in the Available Devices pane. These options are also available in the **File** menu.

When a port refresh is selected, a list of connected devices is displayed in the Connected Devices pane. If problems occur when using the refresh feature while connected to a 4-port replicator, consider restarting the System Maintenance Software to restore the connection.

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Chapter 5

Application and Database Settings

Application Settings

Use the Application Settings dialog box to configure the available application options for the System Maintenance Software. To open the Application Settings dialog box:

1. On the main menu, choose **Options > Application Settings**.
The Application Settings dialog box is displayed.
2. To change an application setting, click the column to the right of that setting and make the appropriate change.
3. Close the Application Settings dialog box to save your changes.

Setting	Description
Flash <ul style="list-style-type: none"> • Allow serial number update • Default SpO₂ Module for Flash 	<p>If true, the device serial number can be set through flash tasks.</p> <p>When flashing the firmware on an SpO₂ Module, either Nellcor® or Masimo® can be selected as the default firmware.</p>
General <ul style="list-style-type: none"> • Beep for user input • Beep on finished task • Enable task list edit button • Show only connected ports 	<p>If true, causes application to beep when a running task requires attention.</p> <p>If true, causes application to beep when a task finishes.</p> <p>If true, task list edit button is enabled, allowing Task List to be edited.</p> <p>If true, only communication ports that have devices connected to them are displayed in the Connected Devices pane. If false, all available ports are displayed.</p>
Log Downloading <ul style="list-style-type: none"> • Clear downloaded CQI logs • Clear downloaded logs 	<p>If either clear downloaded logs setting is set to True, the software automatically clears the log during download. If you are performing an instrument investigation, CareFusion recommends that the logs not be cleared until the investigation is completed.</p> <p>If true, CQI logs are cleared from connected devices after download.</p> <p>If true, all battery, error, and event logs (other than CQI logs) are cleared from connected devices after download. This setting does not affect whether CQI logs are cleared after downloading. The clearing of CQI logs after downloading is managed separately from all other settings used to clear logs.</p>
Package Components <ul style="list-style-type: none"> • Ignore incomplete packages 	<p>If true, any empty packages are ignored and do not produce an error.</p>

Database Settings

Multiple database definitions can be defined for the three System Maintenance Software databases (Maintenance database, Maintenance Log database, and CQI database). Each definition contains the server name and authentication for the databases. Once defined, the user can select which database to use from this list of defined databases.

Once defined, do not change database settings unless you have a valid reason for doing so. For example, you may need to select a site-specific database in an environment that supports multiple hospital sites.

The System Maintenance Software automatically connects to the default database locations, but additional customizing may need to occur if your database definition is different from these default parameters.

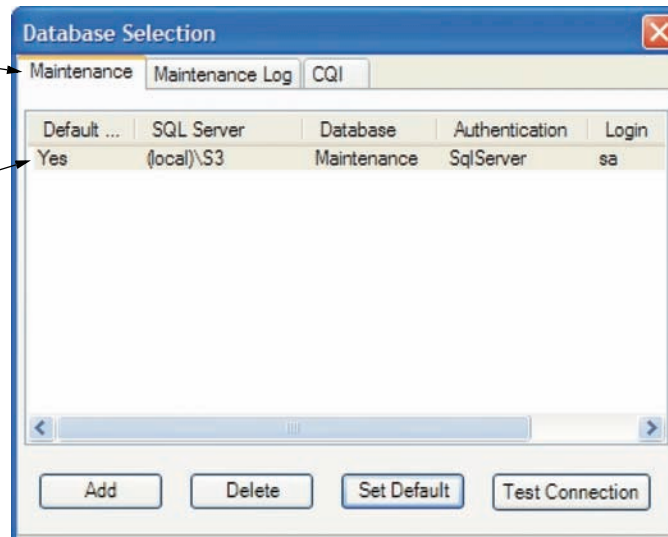
Open Database Settings

On the main menu, choose **Options > Database Settings**.

The Database Selection dialog box is displayed.

Select a tab to display the list of database definitions for the selected System Maintenance Software database.

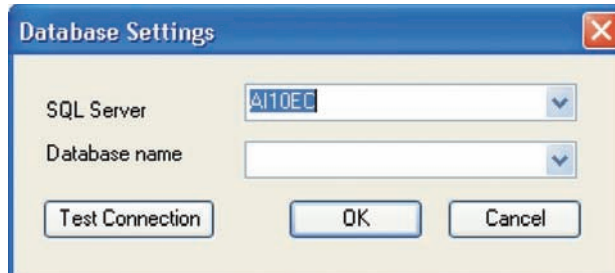
Database definitions relating to the selected tab display here. The active definition has **Default** set to **Yes**.



Add a Database Definition

1. Click the tab of the desired System Maintenance Software database.
2. Click **Add**.

The Database Settings dialog box is displayed.



3. Select an SQL Server.
4. Select a Database name on this SQL Server.
5. Click **Test Connection** to test the connection to the server.
6. Click **OK** to acknowledge the test connection.
7. Click **OK** to save the definition.

Delete a Database Definition

1. Click the tab of the desired System Maintenance Software database.
2. Click the database definition that you want to delete.
3. Click **Delete**.
A confirmation dialog box is displayed.
4. Click **OK** to delete the database definition.

Set Active Database Definition

1. Click the tab of the desired System Maintenance Software database.
2. Click the database definition to make it active.
3. Click **Set Default**.

The System Maintenance Software now uses the selected definition as the active database and begins logging data to it as data is collected.

Test Database Definition Connection

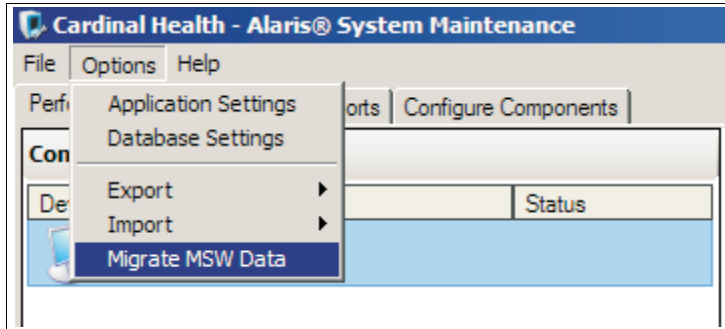
1. Click the tab of the desired System Maintenance Software database.
2. Click the database definition to select it.
3. Click **Test Connection**.

If the connection fails, check the database definition for errors.

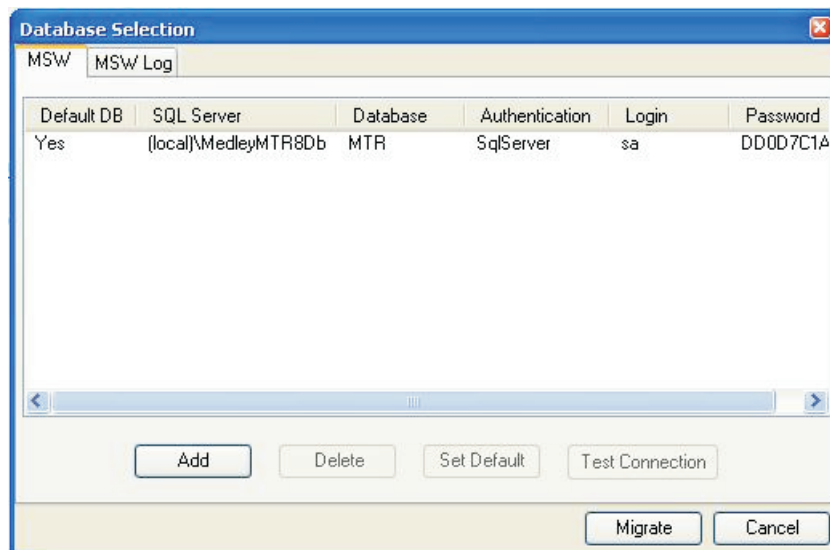
Migrate MSW Databases

There are two separate MSW application databases, MSW and MSW Log. Use the following procedure to migrate the MSW databases to the System Maintenance Software database.

1. On the main menu, click **Options > Migrate MSW Data**.



The Database Selection screen is displayed:



2. If the desired MSW or MSW Log databases are not listed, continue with the following steps; otherwise, proceed to step 3.

NOTE

The desired MSW databases (MSW and MSW Log) must be configured as default DB.

- a. Click the tab for the applicable database and then click **Add**. The Database Settings screen is displayed.



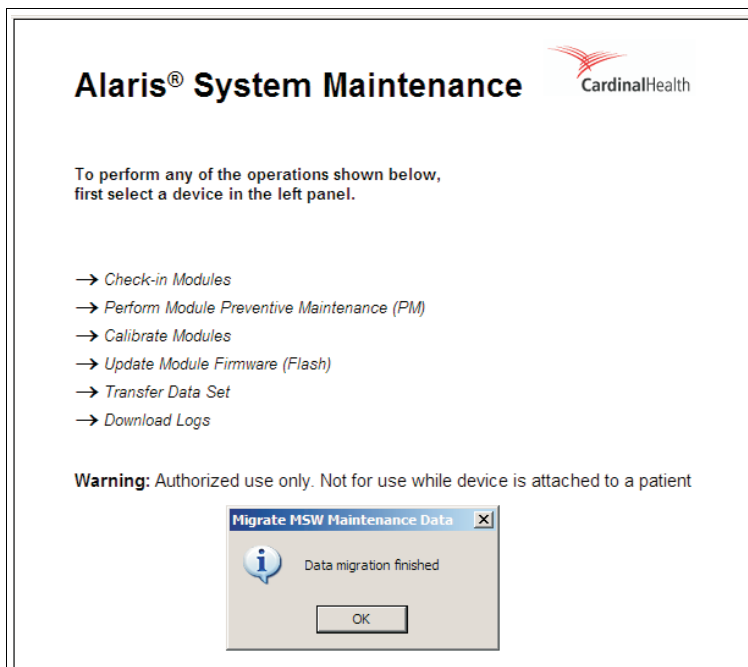
- b. Select the desired server from the **SQL Server** drop-down list.
- c. From the **Database name** drop-down list, select MTR for the MSW database or IMSLog for the MSW Log database.

NOTE

If the required database is not in the **Database name** drop-down list, the selected SQL Server is not correct.

- d. Click **Test Connection** to have the software verify that the MSW database configuration is correct.
3. Click on desired database.
 4. Click **Migrate**—located at the bottom of the Database Selection screen—to start data migration.

The data migration progress screen is displayed. When the migration process is complete, the application displays the status of the migration.



Chapter 6

Preventive Maintenance

Introduction

WARNING

At no time should the System Maintenance Software be used while the Alaris® System is connected to a patient.

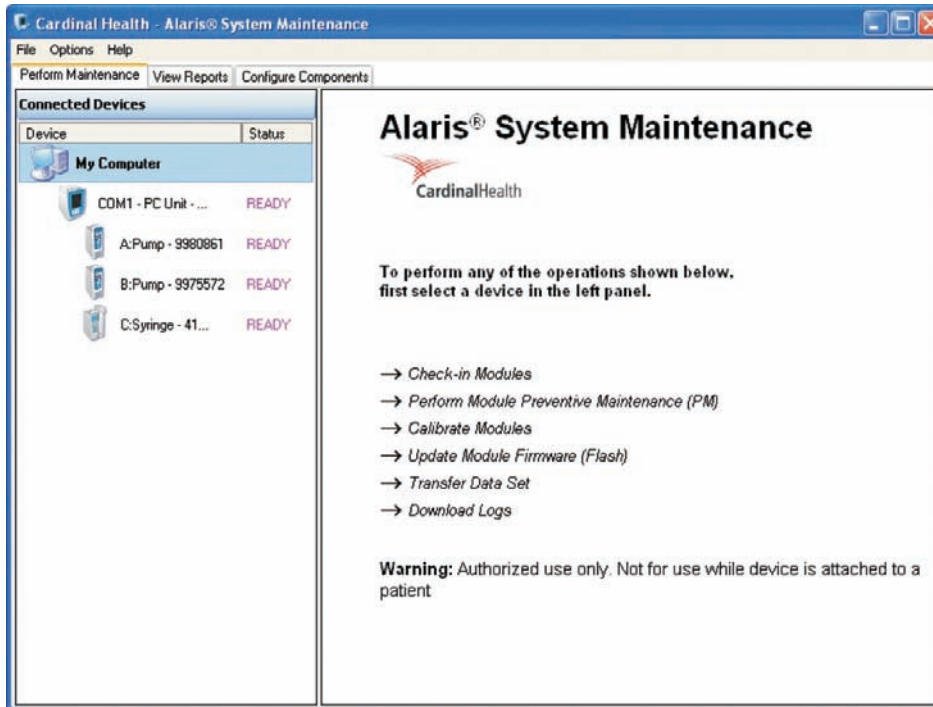
Preventive maintenance can be performed on connected modules by selecting from a list of automated tasks. Tasks can be individual tasks that exist in the list of master tasks or task groups that contain a collection of the individual tasks required to maintain devices.

Standard task groups include:

Check-in	<p>Tasks required to check any module newly received into the facility. Run check-in on a newly received module to confirm that it was not damaged during transport and is ready to be put into use. System Maintenance Software guides you through check-in.</p> <p>The final task in the Check-in task group allows you to set a Preventive Maintenance (PM) Reminder Date to remind the user when preventive maintenance is due.</p>
Preventive Maintenance	<p>Tasks required to perform regular maintenance on a module.</p> <p>Run preventive maintenance tests a minimum of once a year to confirm that modules are performing correctly. System Maintenance Software guides you through preventive maintenance tests.</p> <p>Final task in Preventive Maintenance task group allows you to set the next PM Reminder Date to remind the user when preventive maintenance is due.</p>
Calibration	<p>Tasks required to test whether calibration is required and to calibrate connected modules.</p> <p>The calibration task group applies to EtCO₂ Module, Syringe Module, PCA Module, and Pump Module.</p>

General Steps

The main window is shown below. When you click a node such as **My Computer** in the Connected Devices pane, information on how to proceed is displayed in the right pane.



1. Click the **Perform Maintenance** tab.
2. Select the module (device).

The Tasks list is displayed.

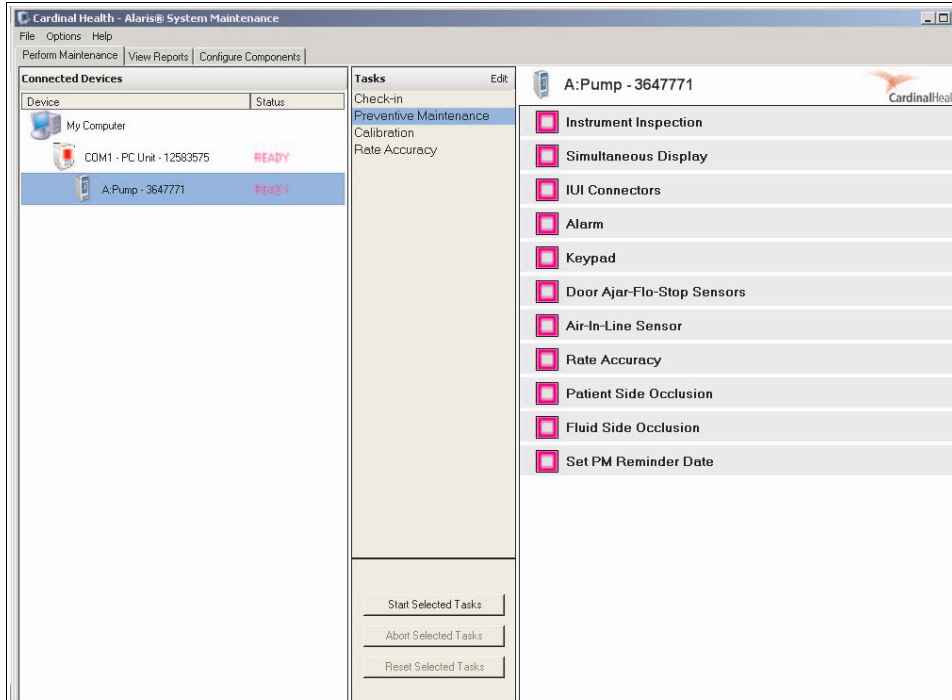
NOTE

- The tasks in the Tasks list vary depending on which module you have selected and how the tasks are configured.
- You can edit the Tasks list, if desired (see *Tasks List* on page 168).

- Click **Preventive Maintenance** in the Tasks list (Pump Module screen used as an example in following illustration).

NOTE

As a shortcut, you can double-click the task to select it and start it immediately.



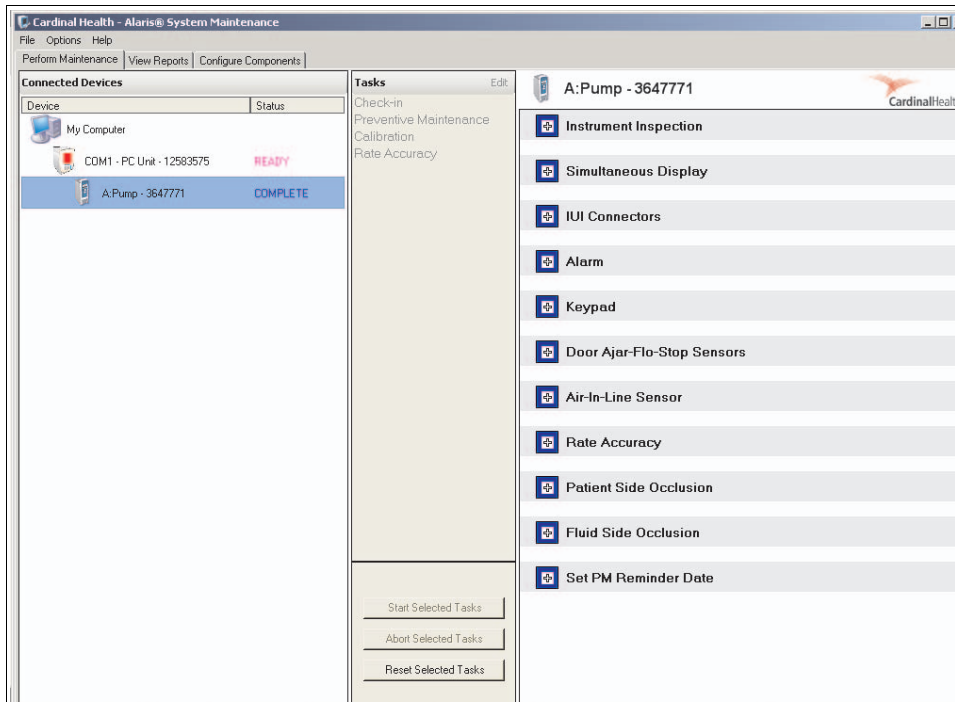
- If Preventive Maintenance was not automatically started, click **Start Selected Tasks**.
 - Follow all on-screen instructions carefully to ensure the successful completion of the task.
- If a task fails, the System Maintenance Software asks you if you want to rerun the failed task.

NOTE

At the end of some tests you are asked to indicate if the test passed or failed. Other tests automatically indicate a pass or fail result.

- To rerun the task, click **Yes**.
- To continue to the next task in a task group, click **No**.
- To correct a failure, abort the selected task and refer to the applicable Technical Service Manual.

6. When finished, click **Reset Selected Tasks** to return a module to the **READY** state (Pump Module screen used as an example in following illustration).

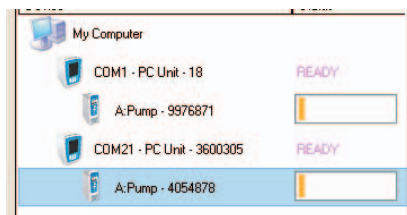


Run/Abort Tasks

Run Tasks Concurrently

You can run selected tasks concurrently on different connected modules. One PC Unit and a serial port is needed for each module being tested. Once you have started selected tasks on one connected device, select other devices in the Connected Devices pane and start tasks for those devices, if desired.

The status bar displays devices that are currently running tasks. If attention is required by a device to continue a task, the device progress bar and the Details Pane Task button for that task will be orange.



Select the device in the Connected Devices pane to display its task instructions and follow the displayed task instructions.

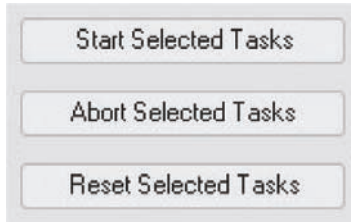
The System Maintenance Software sounds an alarm when a device needs attention or when a task is finished, if these options are set. For information on setting these options, see *Application Settings* on page 30.

Abort Tasks

You can abort (cancel) any running task that is currently selected in the Connected Devices pane.

Click **Abort Selected Tasks**.

The status of the task is reset to **CANCEL**.



Coloring Scheme

The preventive maintenance test results are identified using the following color scheme:

Blue—test passed

Brown—test aborted

Red—test failed

Orange—test is running, but waiting for user input

Purple—test has not yet been run

Green—test is automatically running (not requiring user input at this time)

Example:



Preventive Maintenance

To perform preventive maintenance, see the procedure in this section that applies to the module being tested:

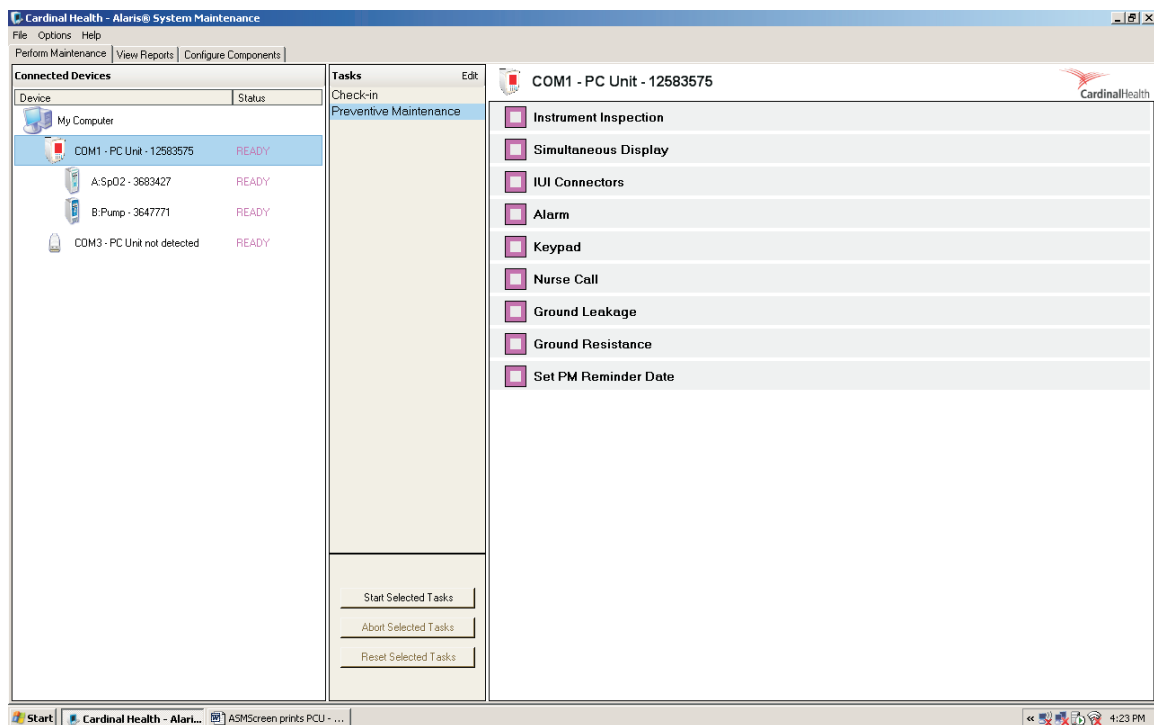
- PC Unit on page 41*
- Pump Module on page 47*
- PCA Module on page 59*
- Syringe Module on page 67*
- SpO₂ Module on page 77*
- EtCO₂ Module on page 84*
- Auto-ID Module and Handheld Scanner on page 91*

PC Unit

Test Equipment

Test Equipment	Manufacturer	Model Number	Application
analyzer, electrical safety	Fluke BioMedical	232D or equivalent	ground tests

1. In the Connected Devices pane, click the PC Unit to be tested.
2. Click **Preventive Maintenance** in the Tasks list (double-click to begin testing immediately).



3. Follow the instructions displayed on the screen for each test.

NOTE

See *PC Unit Tasks* on page 170 for an explanation of each task.

4. Instrument Inspection:

Step 1 Clean any surfaces where solution or obstructions have accumulated.

Examine the instrument for overall condition.

Verify that

1. The case is clean and free from residue, especially near moving parts.
2. Keys, labels, and markings are free of damage and are clearly legible.
3. No tape or other foreign material is on the side of the module.
4. Visually inspect all IUI connectors for damage, bent pins, and signs of corrosion (green material). Replace if found.
5. Case has no apparent damage.
6. IUI latch moves freely.
7. Each key is illuminated by backlight (you may need to cover the keys in order to see the backlight).

Step 2 The pole clamp should be secure and functioning. If the instrument is mounted on a pole or stand, examine the condition of the mount, the pole, and the stand.

Examine the power cord assembly for:

1. Signs of damage, cuts, or deformities in the cord.
2. Integrity of hospital grade power plug. Attempt to wiggle the blades to determine that they are secure.
3. Appropriate tension and connection if the IV pole has electrical receptacles for accessories.
4. Examine the strain reliefs at both ends of the line cord. Be sure they hold the cord securely.

If any damage is suspected, replace the entire cord.

Step 3 Determine Pass/Fail

Press the 'Pass' button if the test has successfully completed. If any errors have occurred, press the 'Fail' button.

If the test failed, click **Abort Selected Tasks**—in the Tasks list—to discontinue testing. Correct the failure(s) before continuing with the tests.

5. Simultaneous Display:

If you want the test to run continuously, click the **Run test in continuous mode** checkbox.

Step 1 This test will light all display elements on the LCD by displaying screens of solid colors.

Current display settings:

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Run test in continuous mode

While the test is running, check to make sure the device displays a series of solid screens, black, gray, white, red, green, and blue. Note that the test may take a few moments to complete after clicking 'Stop'. If running the test in continuous mode, uncheck the continuous mode checkbox or press 'Stop' to stop the test.

When ready to continue, press the 'Next' button.

- Look for missing pixels (display elements) as each screen is displayed.
- The Model 8000 displays dark gray and light gray screens. The Model 8015 displays dark gray, light gray, white, red, green, and blue screens.
- If the screen brightness or contrast appears weak, check the contrast setting (refer to applicable Alaris® System DFU).

Step 2 Determine Pass/Fail

Press the 'Pass' button if the test has successfully completed. If any errors have occurred, press the 'Fail' button.

6. IUI Connectors:

Step 1 Visually inspect all IUI connectors for damage, bent pins, and signs of corrosion (green material). Replace if found.

Connect two modules (not Auto-ID) to the PC Unit (one on each side) 'sandwiching' the PC Unit (if not already done).

Wait for the Channel ID to appear on the attached units before pressing 'Next'.

'Sandwiching'—the PC Unit has one module attached to each side.

Step 2 Result of device poll

Summary of the number of devices detected

Expected number of devices :	3
Actual Number of devices :	3
	Pass

Press 'Finish' to continue

7. Alarm:

Alarm

Step 1 Do you hear the PC Unit speaker alarming?

Yes No

If the audio is weak, check the audio volume setting (refer to applicable Alaris® System DFU).

Step 2 A Pass result will be entered for this device.

Finish

8. Keypad:

NOTE

Press and hold the Tamper Switch (on back of PC Unit) for at least two seconds, until there is a response to the test.

Step 1 Please press every key and verify the key pressed. Pressing 'CANCEL' twice in rapid succession will stop the test. To test the tamper switch hold it in for at least 2 seconds. When the test is completed, or can't be finished for any reason (like a key not registering), press the 'Next' button.

S1			S6
S2			S7
S3	TAMPER SWITCH		S8
S4			S9
S5			S10
SILENCE			
OPTIONS			
S11	S12	S13	S14
1	2	3	ARROW UP
4	5	6	ARROW DOWN
7	8	9	ENTER
CLEAR	0	DECIMAL POINT	CANCEL

Next

Press each key one time.

- Green text—the key was pressed one time and passed the test.
- Red text—the key was pressed two times, or it was pressed one time and failed the test. If the text is red because you pressed the key more than one time, you can indicate that the test has passed.

Step 2 Test has passed, press the 'Finish' button

Finish

9. Nurse Call:

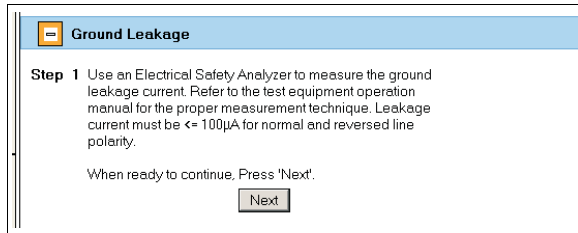
This test is performed only if a Nurse Call is installed and applies only to the Model 8000.

Nurse Call

Step 1 Nurse call is not installed.

10. Ground Leakage:

The ground leakage test is not performed by the System Maintenance Software. The leakage current table provided below reflects IEC/UL 60601-1 requirements. The limits identified in the table are to be used in place of the 100 μ A leakage current limit specified by the software.

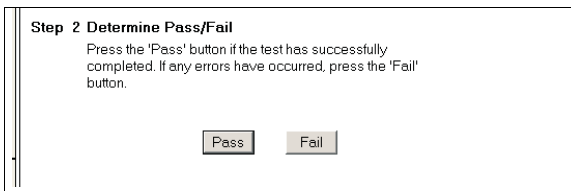


Use a Fluke Biomedical Model 232D or equivalent electrical safety analyzer to measure the ground leakage current. Refer to the test equipment's operation manual for the proper measurement technique.

The line voltage must be the same as the voltage that powers the tester and then passes to the instrument being tested.

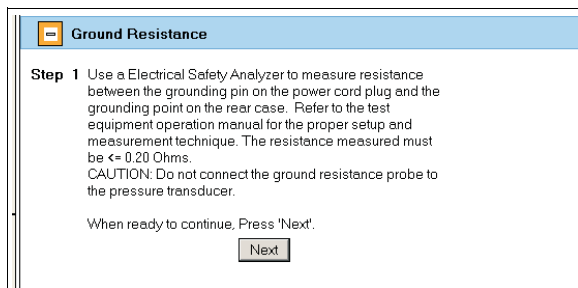
Use the following table—in place of the requirements identified by the software—to determine the leakage current requirements for normal and reversed line polarity. The limits are 100 μ A below the IEC/UL 60601-1 standard limits to account for test variances.

	AC Supply	Leakage Current—Maximum
Japan	100 VAC	400 μ A
North America	120 VAC	200 μ A
Rest of World	230 VAC	400 μ A
Australia New Zealand	240 VAC	400 μ A



11. Ground Resistance:

The ground resistance test is not performed by the System Maintenance Software. Use the following instructions and illustration to perform this test—in place of the instructions displayed by the software.



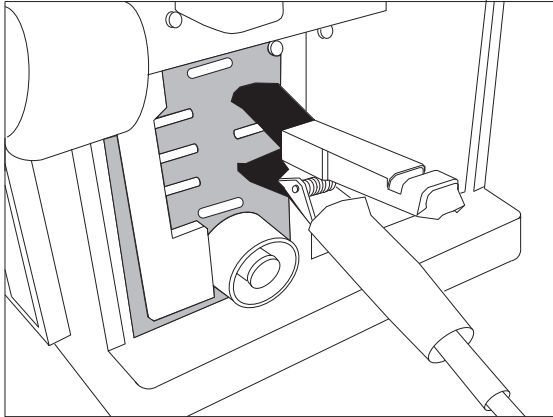
Use a Fluke Biomedical Model 232D, an Ohm Meter, or an equivalent electrical safety analyzer to measure the ground resistance between the ground pin on power cord plug or power cord receptacle, and the rear panel of the PC Unit. Refer to the test equipment's operation manual for the proper measurement technique.

The maximum allowable resistance is 1 Ohm.

The following illustration is a partial view of the rear of the PC Unit. In the illustrated example of a ground contact, a Kelvin Clip is clipped to a speaker slot on the rear panel. The only acceptable ground contact area is the rear panel—gold anodized color metal (shaded area in illustration).

NOTE

The following illustration reflects the use of the Kelvin Clip that is provided with the Fluke tester. If another tester or Ohm Meter is used, the test measures the resistance between the rear panel (shaded area in illustration) and the ground pin on the power cord plug or power cord receptacle.



Step 2 Determine Pass/Fail

Press the 'Pass' button if the test has successfully completed. If any errors have occurred, press the 'Fail' button.

Pass Fail

12. Set PM Reminder:

- If **Enable Maintenance Reminder** is selected, a Data Set is loaded and the configuration option for the Data Set is set to **Disable**. The Data Set setting overrides the System Maintenance Software selection.
- The next maintenance reminder date defaults to 12 months. The reminder can be set to an earlier date but it cannot be set to a date past the 12-month default.

Set PM Reminder Date

Step 1 Next maintenance: 10/ 2/2010

Enable Maintenance Reminder
 Note: This selection will be overridden if the module has a data set loaded.

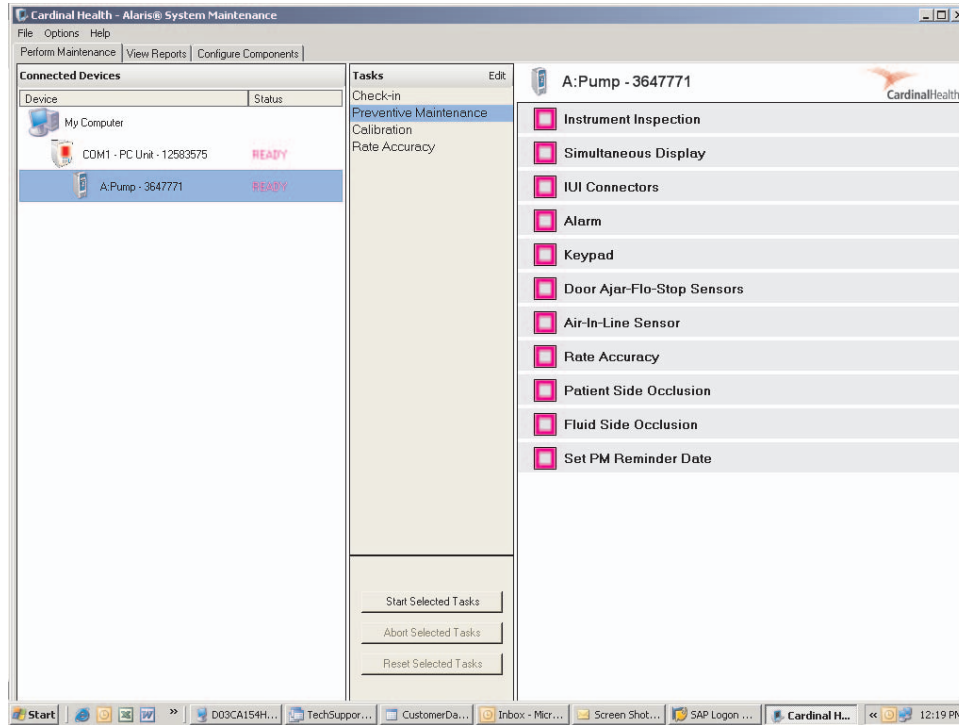
Finish

Pump Module

Test Equipment

Test Equipment	Manufacturer	Model/Part Number	Application
gauge, pressure, digital (peak hold)	Either of the following: <ul style="list-style-type: none"> • Heise (www.heise.com) • Ashcroft (www.ashcroft.com) 	<ul style="list-style-type: none"> • PTE-1 (accuracy from 0.1 to 0.025% span) • 2089, 2086, or 2084 (accuracy from $\pm 0.05\%$, 0.10%, or 0.25% of span) • or an equivalent gauge with: <ul style="list-style-type: none"> (a) unit of measure in mmHg (b) accuracy of $\pm 1\%$ (c) range of 0-1500 mmHg 	pressure test
hemostat	N/A	N/A	fluid-side occlusion test
IV pole, standard	CareFusion	903-0336 or equivalent	rate, pressure, and occlusion tests
IV sets <ul style="list-style-type: none"> • IV set, standard, without check valve • IV set, calibration • IV set, calibration 	<ul style="list-style-type: none"> • CareFusion • CareFusion • CareFusion 	<ul style="list-style-type: none"> • 2210-0500 • 8100-RCS • 8100-PCS 	rate and pressure tests
IV solution container (bag preferred)	N/A	N/A	rate and pressure tests
luer lock, female, 1/8"-27 NPT	Cole-Parmer (www.coleparmer.com)	K-45503-78 or equivalent	pressure test
reducer, female, 1/4" NPT (F) x 1/8" NPT (F)	Cole-Parmer (www.coleparmer.com)	K-06349-91 or equivalent	pressure test
scale, digital	Acculab (www.accumlab.com)	VIC-212 (VICON Series) or equivalent with accuracy of $\pm 0.01\text{g}$	rate and pressure tests
tubing, silicone	CareFusion	303109 or equivalent	pressure test
T-fitting	CareFusion	303815 or equivalent	pressure test
valve, 3-way	CareFusion	97555 or equivalent	pressure test

1. In the Connected Devices pane, click the Pump Module to be tested.
2. Click **Preventive Maintenance** in the Tasks list (double-click to begin testing immediately).



3. Follow the instructions displayed on the screen for each test.

NOTE

See *Pump Module Tasks* on page 175 for an explanation of each task.

4. Instrument Inspection:

Step 1 Clean any surfaces where solution or obstructions have accumulated.

Examine the instrument for overall condition.

Verify that

1. The case is clean and free from residue, especially near moving parts.
2. Keys, labels, and markings are free of damage and are clearly legible.
3. No tape or other foreign material is on the side of the module.
4. Visually inspect all IUI connectors for damage, bent pins, and signs of corrosion (green material). Replace if found.
5. Case has no apparent damage.
6. IUI latch moves freely.
7. Each key is illuminated by backlight (you may need to cover the keys in order to see the backlight).

Step 2 Interior Door Inspection

Open door and verify:

1. Door Latch Spring Test: The latch stays up and does not sag.
2. Sear Spring Test: When the sear is depressed and released, it springs back into place.
3. Platen Spring Test: The door/platen moves freely and stays open when the door is opened.
4. The mechanism seal is not torn or worn.

After the inspection is complete shut the door.

Step 3 Determine Pass/Fail

Press the 'Pass' button if the test has successfully completed. If any errors have occurred, press the 'Fail' button.

If the test failed, click **Abort Selected Tasks** to discontinue testing. Correct the failure(s) before continuing with the tests.

5. Simultaneous Display:

If you want the test to run continuously, select **Run test in continuous mode**.

Simultaneous Display

Step 1 Run test in continuous mode

Verify:

- 1) Channel ID display.
- 2) Scrolling display.
- 3) Rate display
- 4) Lighthouse LED's
- 5) No missing pixels

Note that the test may take a few moments to complete after clicking 'Stop'. If running the test in continuous mode, uncheck the continuous mode checkbox or press 'Stop' to stop the test.

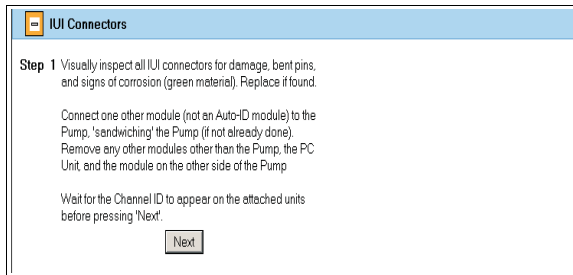
When ready to continue, press the 'Next' button.

- **Channel ID display**—all segments must light.
- **Scrolling display** and **Rate display**—must flash dashes and zeros.
- **Lighthouse LED's**—all LEDs must light.

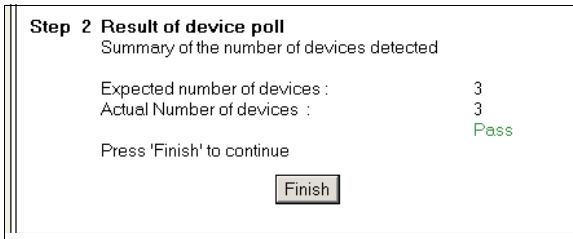
Step 2 Determine Pass/Fail

Press the 'Pass' button if the test has successfully completed. If any errors have occurred, press the 'Fail' button.

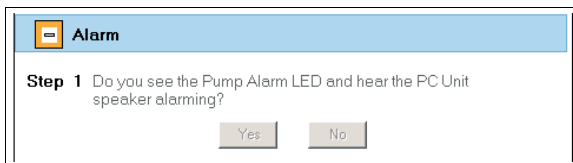
6. IUI Connectors:



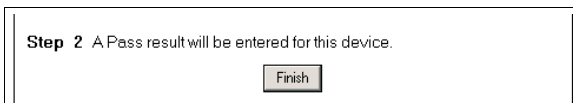
'Sandwiching'—the module being tested is connected between the PC Unit and another module.



7. Alarms:

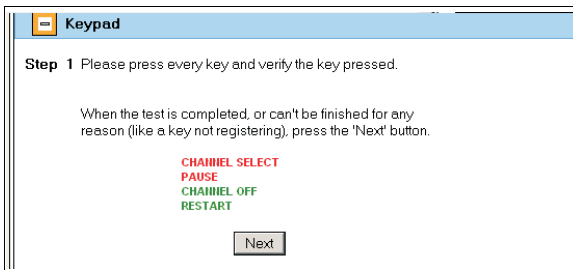


If the audio is weak, check the audio volume setting (refer to applicable Alaris® System DFU).



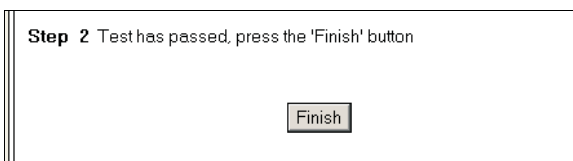
If the audio is weak, check the audio volume setting (refer to applicable Alaris® System DFU).

8. Keypad:



Press each key one time.

- Green text—the key was pressed one time and passed the test.
- Red text—the key was pressed two times, or it was pressed one time and failed the test. If the text is red because you pressed the key more than one time, you can indicate that the test has passed.



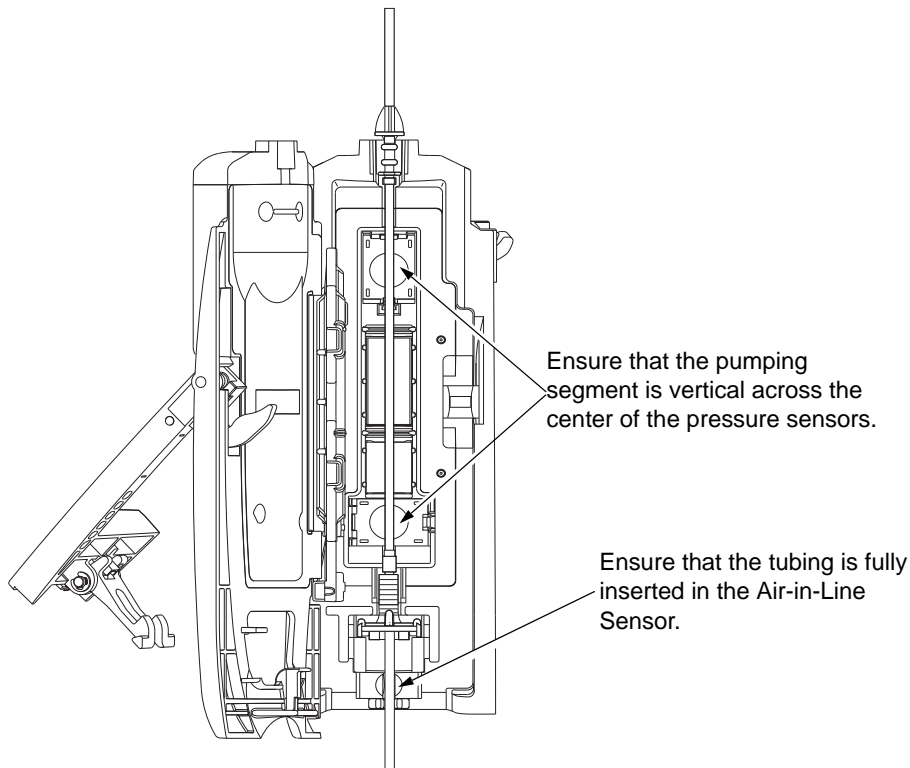
9. Door Ajar-Flo-Stop Sensors:

Door Ajar-Flo-Stop Sensors	
Step 1 Open the door to the Pump Module to verify the correct operation for Door Ajar and Flo-Stop sensors.	
<input type="button" value="Next"/>	

Step 2	Expected Door Open Result:	True
	Actual Door Open Result:	True
	Expected Flow Stop Open Result:	False
	Actual Flow Stop Open Result:	False
	Test Result:	Pass
Step 3 Load a dry set into the Pump Module and close the door to verify the correct operation for the Door Ajar and Flo-Stop sensors.		
<input type="button" value="Next"/>		

Load Administration Set

Install an administration set as shown for any Pump Module test. Unless otherwise specified, in order to run a Pump Module test, the set must be loaded and the Pump Module door must be closed.



Step 4	Expected Door Open Result:	False
	Actual Door Open Result:	False
	Expected Flow Stop Open:	True
	Actual Flow Stop Open:	True
	Test Result:	Pass
Step 5 A Pass result will be entered for this device.		
<input type="button" value="Finish"/>		

10. Air-in-Line Sensor:

■ **Air-In-Line Sensor**

Step 1 Load an empty dry set into the Pump Module and close the door.

Step 2

Expected Sensor Value:	True
Actual Sensor Value:	True
Results of Sensor Test:	Pass

Step 3 A Pass result will be entered for this device.

11. Rate Accuracy:

CAUTION

If the Pump Module fails the rate accuracy test, calibration is required. Failure to perform calibration tests might result in improper operation.

Set up the Pump Module for the Rate Accuracy test.

Rate Accuracy Test Setup**CAUTION**

- If the Pump Module fails the rate accuracy verification test, calibration is required. Failure to perform calibration tests might result in improper operation.
- To ensure accurate rate calibration, use only rate calibration sets. Rate accuracy calibration sets are valid for 60 calibrations and must be replaced after 60 uses.
- Use distilled water at room temperature of 41°F to 104°F (5°C to 40°C). If the water temperature is not within this range, the readings might be inaccurate.

Use this setup to perform the following tests:

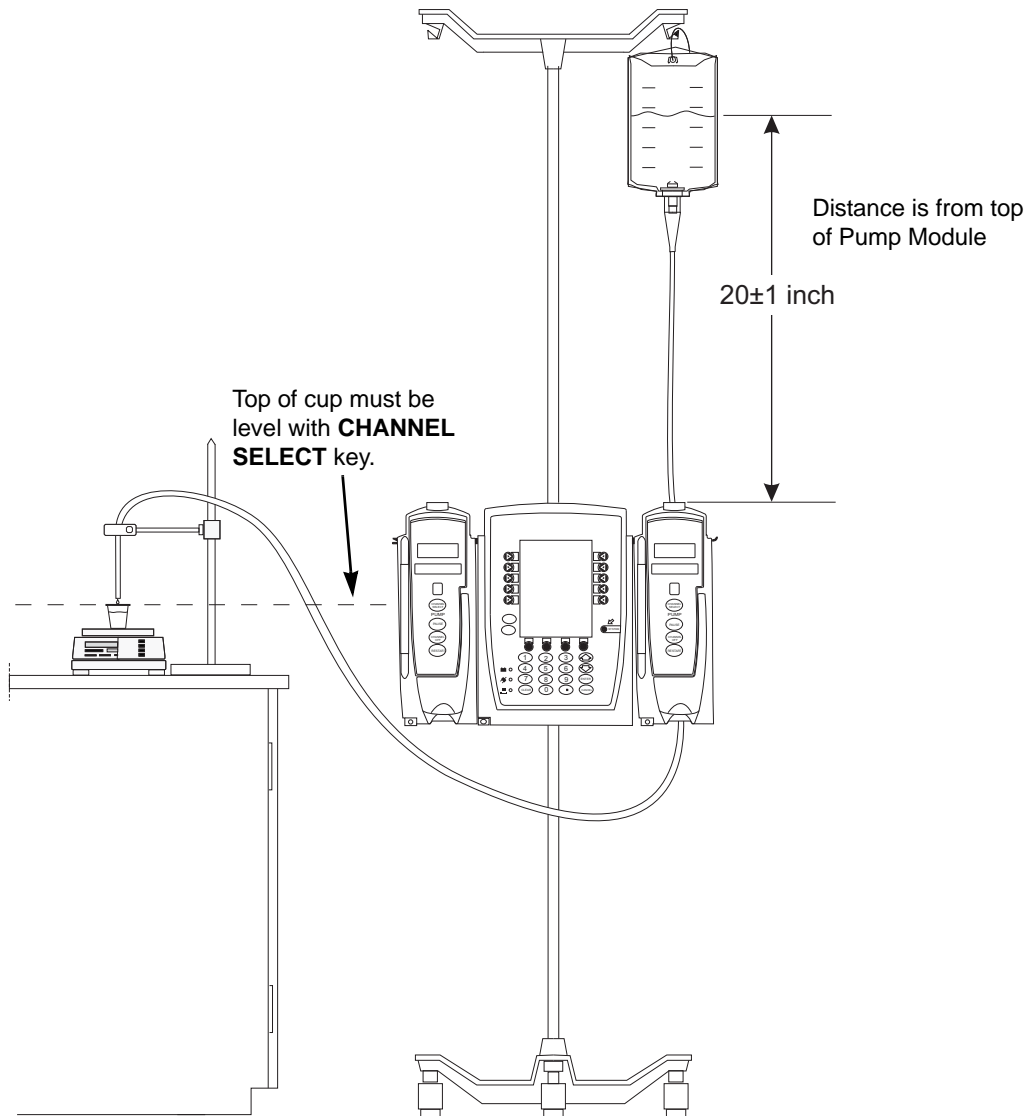
- Rate accuracy verification—Tasks: Rate Accuracy, Check-in, and Preventive Maintenance.
- Rate accuracy calibration pre-test—Task: Rate Accuracy Pre-Test.
- Rate accuracy calibration—Task: Rate Calibration.
- Rate accuracy calibration post-test—Task: Rate Accuracy Post-Test.

All tests use a standard set except for the rate accuracy verification (Task: Preventive Maintenance) and the rate accuracy calibration. These tests require the use of the Model 8100-RCS rate calibration set.

Verify that the scale is calibrated according to the manufacturer's instructions. Prime the fluid lines and keep them free of air bubbles while running the test.

If the rate accuracy calibration pre-test fails, perform the rate accuracy calibration. Following calibration, run the rate accuracy calibration post-test using a different rate calibration set to test the module with the newly calibrated parameters.

—See the test setup illustration on the following page.—



Rate Accuracy	
Step 1	Load a characterized set and prime it with distilled water. Enter the expected volume provided by the characterization label on the packaging of the set. Passing values are 11.640 to 12.360 mL.
Expected Volume:	<input type="text" value="12.00"/> grams
	<input type="button" value="Next"/>

Ensure that the set is properly loaded (see *Load Administration Set* on page 51). Ensure that there are no air bubbles in the set when priming is completed.

The **Expected Volume** is identified on the label attached to the characterized set.

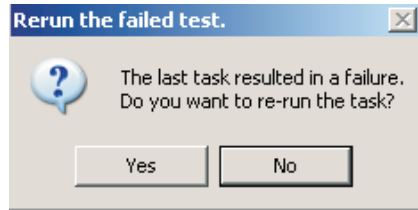
Error message—first occurrence

If an error message appears after clicking **Next**, perform the following steps.

NOTE

If this is not the first occurrence of an error message and the following procedure has already been performed, perform *Error message—repeat occurrence*.

- a. Click **OK** and visually confirm that:
 - The set is properly loaded in the Pump Module.
 - There are no air bubbles in the set.
- b. Click **Yes** on the message that appears.



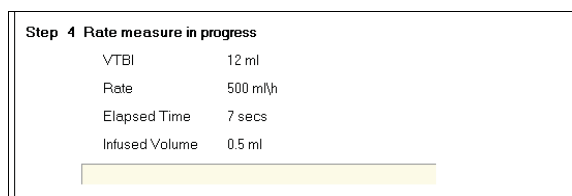
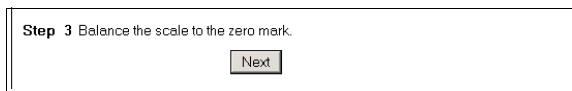
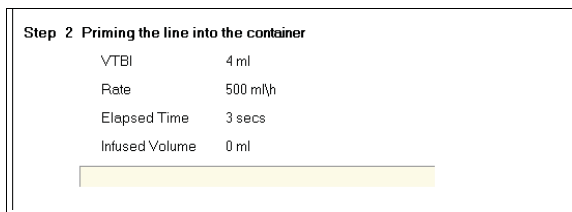
- c. Detach and reattach the Pump Module with set installed.
- d. Repeat the rate accuracy test after the Pump Module reinitializes.

Error message—repeat occurrence

If an error message appears after clicking **Next**, manually connect to the Maintenance Mode and retest, as follows:

- a. Abort the selected task.
- a. Exit the System Maintenance Software.
- b. Turn off the PC Unit.
- c. Ensure that the serial cable is connected between the communications port on the PC Unit and the computer.
- d. Press and hold the Tamper Switch and the **System On** key on the PC Unit until the PC Unit beeps once and displays the **Maintenance Mode** screen.
- e. Press the **Proceed** soft key and then press the **External Communications** soft key.
- f. Start the System Maintenance Software.
- g. Perform Preventive Maintenance.

When performing the Rate Accuracy test, follow the instructions to ensure that the set is properly loaded and that there are no air bubbles in the set when priming is completed.



Ensure that the distilled water is dripping into the cup and that the tubing is not resting on the scale.

Step 5 Enter actual weight reading from scale. Passing values are 11.59 to 12.41 grams.

Actual weight : grams

Step 6 Summary of Test Results

Pump Module Test Result:	Passed
VPMR:	175.3
VTBI:	12 ml
Rate:	500 ml/h
Expected Weight	12.000 grams
Actual Weight	12.0 grams
Acceptable Error	+/-3.400%
Actual Error	0%

12. Patient-Side Occlusion:

Set up the Pump Module for the Patient-Side Occlusion test.

Patient-Side Occlusion Pressure Test Setup

CAUTION

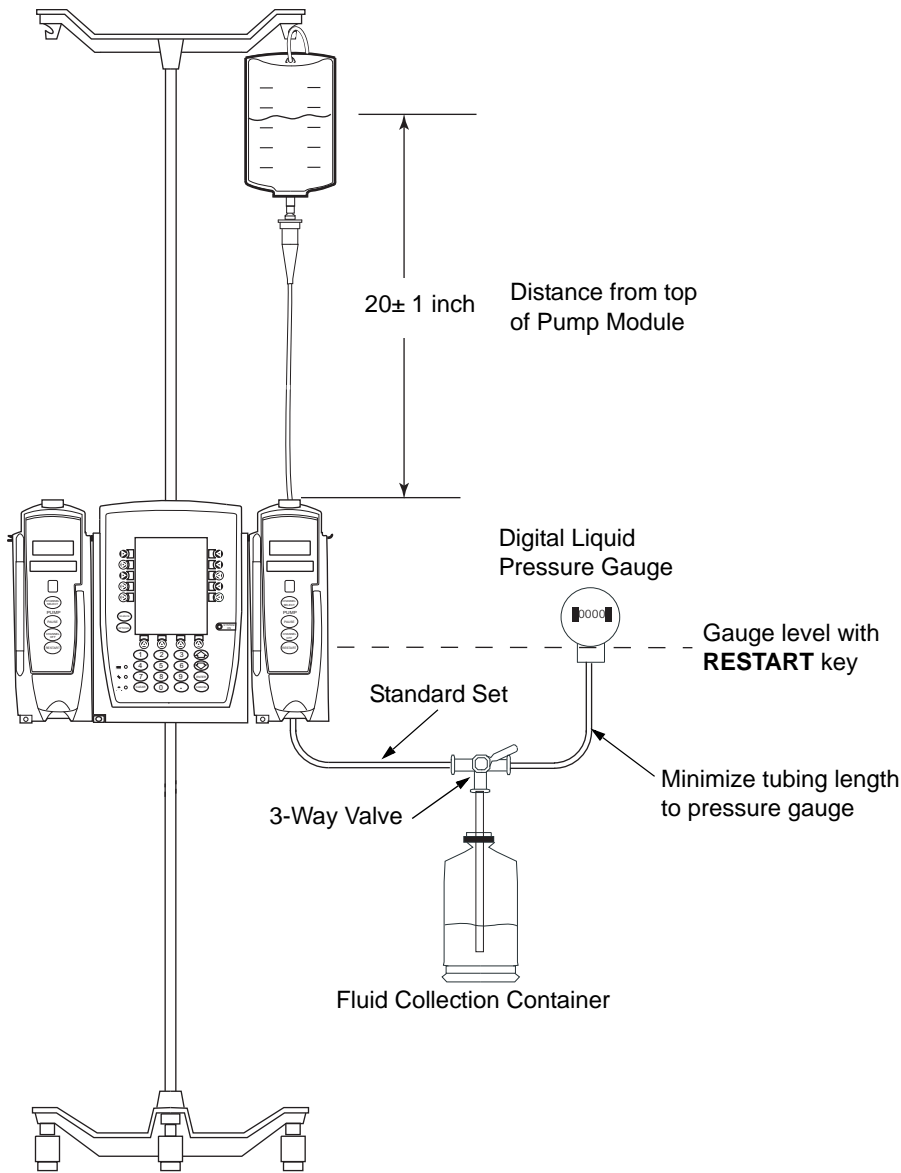
If a Pump Module fails the patient-side occlusion pressure test, calibration is required. **Failure to calibrate and retest** the instrument might result in improper instrument operation.

Use this setup to perform the following tests:

- Patient-side occlusion pressure test—Tasks: Patient-Side Occlusion, Check-in, and Preventive Maintenance.
- Occlusion pressure calibration pre-test—Task: Pressure Pre-Test. If this test fails, perform calibration—Task: Pressure Calibration. After calibration, run the calibration post-test—Task: Pressure Post-Test—with the newly calibrated parameters using the same standard set that was used to perform the pre-test.
- Occlusion pressure calibration post-test—Task: Pressure Post-Test.

All tests use a standard set.

—See the test setup illustration on the following page.—



Patient Side Occlusion

Step 1 Load a primed regular set and close the door. Ensure that the pressure gauge is set to zero.

Step 2 Wait until 0.4 ml has been infused before causing a Patient Side Occlusion.

VTBI	4 ml
Rate	125 ml/h
Elapsed Time	2 secs
Infused Volume	0 ml

After 0.4 ml has been delivered, simulate a patient-side occlusion by flipping the stop cock on the 3-way valve so that fluid is pumping toward the pressure transducer.

Step 3 Pump Occluded. Enter the pressure reading at occlusion.

Pressure reading : psi

Enter the pressure reading at the time of Occlusion.
Passing values are 7.7 to 12.7 psi.

Step 4 A Pass result will be recorded for this device.

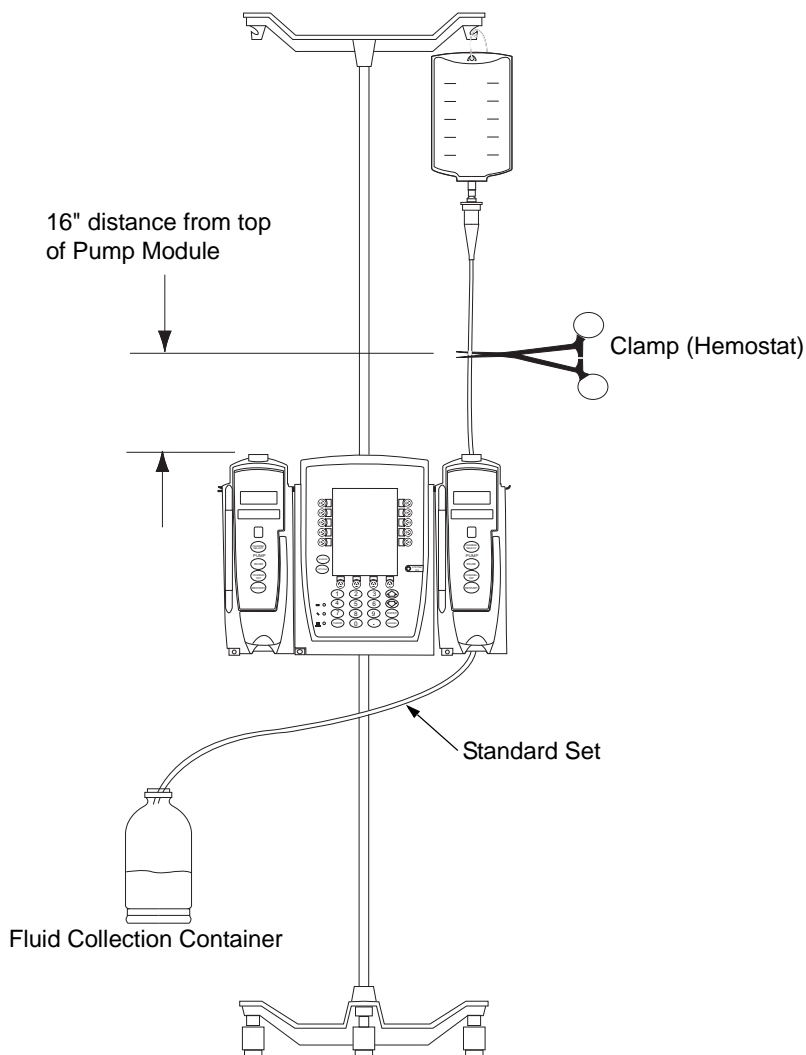
13. Fluid-Side Occlusion:

Set up the Pump Module for the Fluid-Side Occlusion test.

Fluid-Side Occlusion Test Setup

CAUTION

If the Pump Module fails the fluid-side occlusion test, the module is in need of repair.



Fluid Side Occlusion

Step 1 Load a primed regular set into pump module and close door.

Step 2 Wait until 0.4 ml has been infused before causing a Fluid Side Occlusion.

VTBI	4 ml
Rate	125 ml/h
Elepsed Time	5 secs
Infused Volume	0 ml

To simulate a fluid-side occlusion after 0.4 mL has been delivered, clamp the tubing 16 inches above Pump Module.

Step 3 Pump occluded. A Pass result will be recorded for this device.

14. Set PM Reminder:

The next maintenance reminder date defaults to 12 months. The reminder can be set to an earlier date but it cannot be set to a date past the 12-month default.

Set PM Reminder Date

Step 1 Next maintenance:

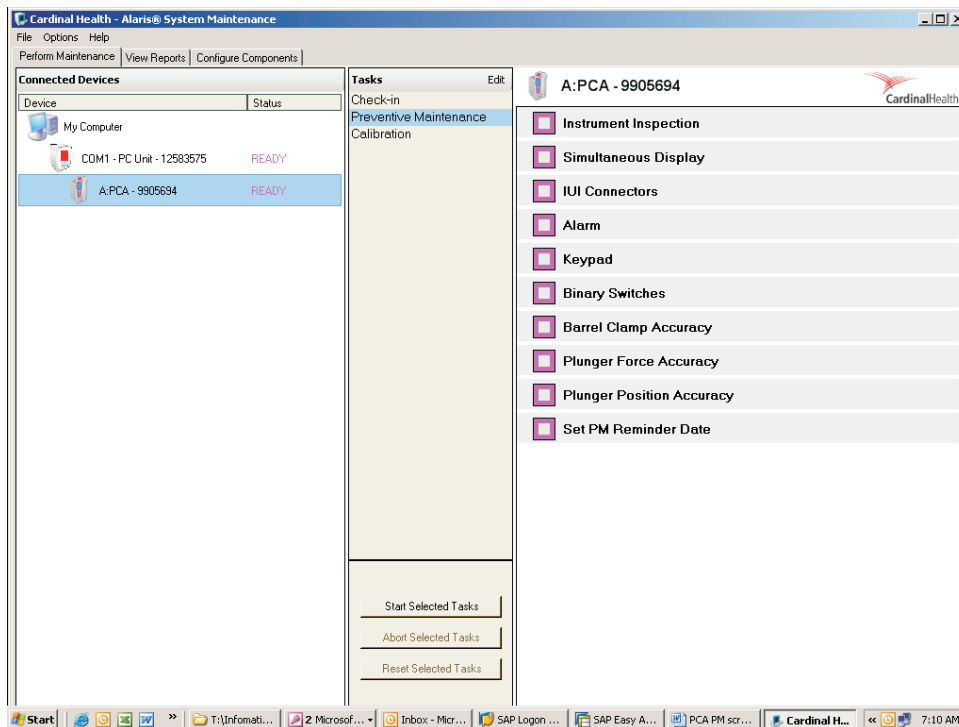
PCA Module

Test Equipment

Test Equipment	Manufacturer	Model/Part Number	Application
door key	CareFusion	10010964	lock and unlock PCA Module door.
Dose Request Cord	CareFusion	10013795	display test
gauge, syringe height	CareFusion	148181-100	plunger position calibration / verification
kit, calibration tools ^a	CareFusion	10010692	all tools required for calibration and verification
kit, force sensor	CareFusion	10010691	force sensor calibration / verification
kit, syringe sizer and height gauge	CareFusion	148182-100	plunger position / barrel size calibration / verification
kit, syringe sizer gauge	CareFusion	148180-100	barrel size calibration / verification

a. To check the spring cartridge verification, preload the cartridge with 0.50 ± 0.05 lbs. Measure the deflection from the preload point after adding 19.5 lbs additional load (20 ± 0.05 lbs total). Acceptable deflection is within 0.891-0.940 inches. Use of Instron or equivalent is recommended.

1. In the Connected Devices pane, click the PCA Module to be tested.
2. Click **Preventive Maintenance** in the Tasks list (double-click to begin testing immediately).



3. Follow the instructions displayed on the screen for each test.

NOTE

See *Syringe Module and PCA Module Tasks* on page 178 for an explanation of each task.

4. Instrument Inspection:

Step 1 Clean any surfaces where solution or obstructions have accumulated.

Examine the instrument for overall condition.

Verify that:

1. The case is clean and free from residue, especially near moving parts.
2. Keys, labels, and markings are free of damage and are clearly legible.
3. No tape or other foreign material is on the side of the module.
4. Visually inspect all IUI connectors for damage, bent pins, and signs of corrosion (green material). Replace if found.
5. Case has no apparent damage.
6. IUI latch moves freely.
7. Each key is illuminated by backlight (you may need to cover the keys in order to see the backlight).

Step 2 Examine the instrument for overall condition.

Verify that:

1. The knob opens and closes freely and the claws open and close with it.
2. The plunger head moves up and down freely when the knob is open.
3. The knob, claws, plunger detect and force sensors have no visible damage.
4. The flange and flange sensor have no visible damage.
5. The barrel clamp has no visible damage and moves/rotates freely.
6. The key lock operates at all three detente positions.
7. The door hinge moves freely and without binding.
8. With the key in the unlock position, the latch does not wiggle freely.
9. With the door closed, and the key locked, rotating the key to the unlocked position causes the door to pop open.
10. The dose request button lights up when pressed.
11. No other external damage is apparent.

Step 3 Determine Pass/Fail

Press the 'Pass' button if the test has successfully completed. If any errors have occurred, press the 'Fail' button.

If the test failed, click **Abort Selected Tasks** to discontinue testing. Correct the failure(s) before continuing with the tests.

5. Simultaneous Display:

If you want the test to run continuously, click the **Run test in continuous mode** checkbox.

Step 1 Run test in continuous mode

Verify:

- 1) Channel ID display.
- 2) Scrolling display.
- 3) Rate display
- 4) Lighthouse LED's
- 5) Dose request cord LED
- 6) No missing pixels

Note that the test may take a few moments to complete after clicking 'Stop'. If running the test in continuous mode, uncheck the continuous mode checkbox or press 'Stop' to stop the test.

When ready to continue, press the 'Next' button.

- **Channel ID display**—all segments must light.
- **Scrolling display** and **Rate display**—must flash dashes and zeros.
- **Lighthouse LED's**—all LEDs must light.
- **Dose request cord LED**—plug dose request cord into Auto-ID Module for this test.

Step 2 Determine Pass/Fail

Press the 'Pass' button if the test has successfully completed. If any errors have occurred, press the 'Fail' button.

6. IUI Connectors:

IUI Connectors

Step 1 Visually inspect all IUI connectors for damage, bent pins, and signs of corrosion (green material). Replace if found.

Connect one other module (not an Auto-ID module) to the PCA, 'sandwiching' the PCA (if not already done). Remove any other modules other than the PCA, the PC Unit, and the module on the other side of the PCA

Wait for the Channel ID to appear on the attached units before pressing 'Next'.

'Sandwiching'—the module being tested is connected between the PC Unit and another module. The other module cannot be a PCA Module.

Step 2 Result of device poll

Summary of the number of devices detected

Expected number of devices :	3
Actual Number of devices :	3
	Pass

Press 'Finish' to continue

7. Alarm:

Alarm

Step 1 Do you see the PCA Alarm LED and hear the PC Unit speaker alarming?

Yes No

If the audio is weak, check the audio volume setting (refer to applicable Alaris® System DFU).

Step 2 A Pass result will be entered for this device.

Finish

8. Keypad:

Keypad

Step 1 Please press every key and verify the key pressed.

When the test is completed, or can't be finished for any reason (like a key not registering), press the 'Next' button.

CHANNEL SELECT
PAUSE
CHANNEL OFF
RESTART

Next

Press each key one time.

- Green text—the key was pressed one time and passed the test.
- Red text—the key was pressed two times, or it was pressed one time and failed the test. If the text is red because you pressed the key more than one time, you can indicate that the test has passed.

Step 2 Test has passed, press the 'Finish' button

Finish

9. Binary Switches:

Binary Switches

Step 1 Please pull and turn the **barrel clamp** into the 'open' clamp position and press 'Next'.

Next

Step 2 Barrel Clamp open position : **Pass**

Please return the **barrel clamp** to the normal position and press 'Next'

Next

Step 3 Barrel Clamp normal position: **Pass**

Please insert a **dose request cord** firmly, depress the **dose request button**, and while keeping the **dose request button** depressed, press 'Next'.

Next

Step 4 Dose request cord in: **Pass**
 Dose request button pressed: **Pass**

Please release the **dose request button** and press 'Next'

Step 5 Dose request button released: **Pass**

Please remove the **dose request cord** and press 'Next'

Step 6 Dose request cord out: **Pass**

Please close and lock the **door** and press 'Next'

The door key is needed to lock the door.

Step 7 Door closed: **Pass**
 Key locked: **Pass**

Please unlock and open the **door** and press 'Next'

Step 8 Door open: **Pass**
 Key unlocked: **Pass**

Please press the **flange detect** switch and press 'Next' while holding it in.

Step 9 Flange Detect in: **Pass**

Please release the **flange detect** switch and press 'Next'

Step 10 Flange Detect out: **Pass**

Please open the claws with the **knob** and press 'Next' while holding them open.

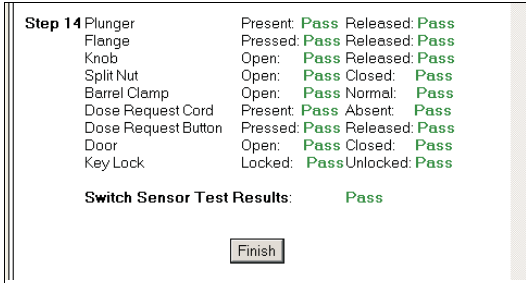
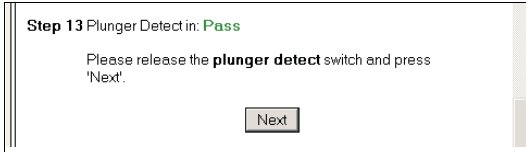
Step 11 Knob open: **Pass**
 Split Nut open: **Pass**

Please release the **knob** allowing the claws to close and press 'Next'.

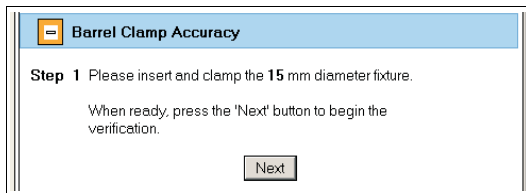
Step 12 Knob closed: **Pass**
 Split Nut closed: **Pass**

Please press the **plunger detect** switch and press 'Next' while holding it in.

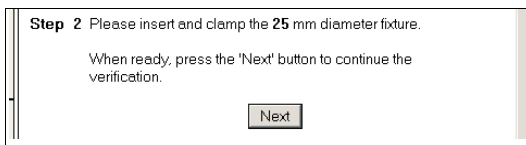
To access the plunger detect switch, open the knob and push the plunger detect switch up.



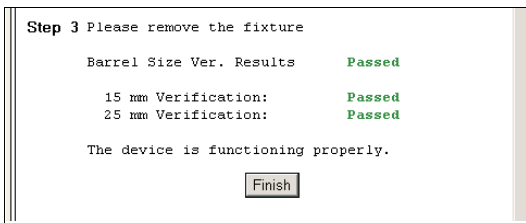
10. Barrel Clamp Accuracy:



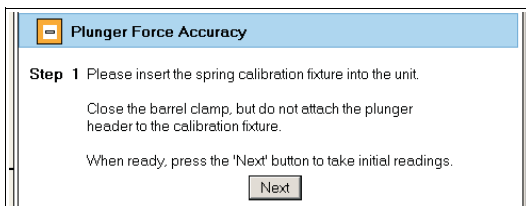
Insert the 15 mm diameter fixture behind the barrel clamp and close the barrel clamp on the fixture.



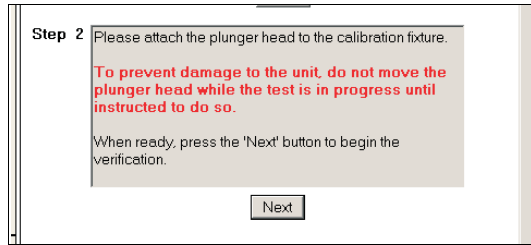
Remove the 15 mm diameter fixture, insert the 25 mm diameter fixture, and close the barrel clamp on the fixture. A popup alerts you if the correct size fixture is not installed.



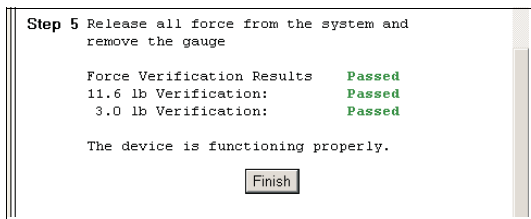
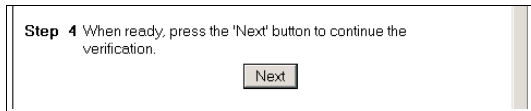
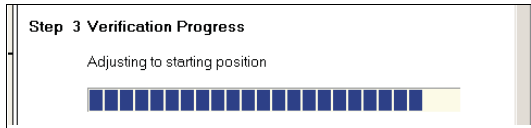
11. Plunger Force Accuracy:



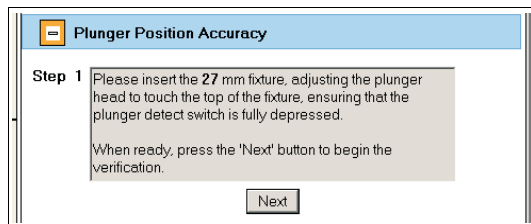
IMPORTANT: To avoid damage to the plunger header, ensure that the plunger header is not attached to the calibration fixture.



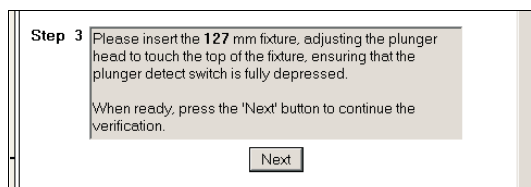
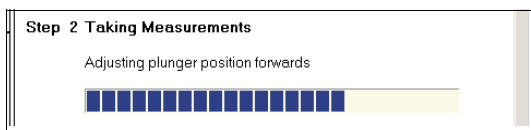
IMPORTANT: To avoid instrument damage, do not move the plunger head during this test.



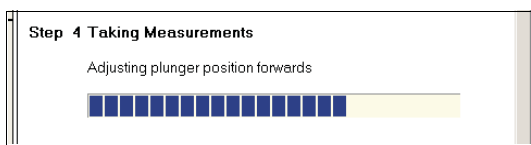
12. Plunger Position Accuracy:

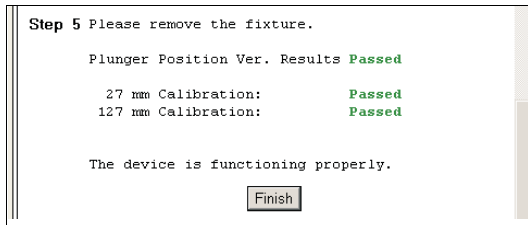


Follow the instructions on the screen and then close the barrel clamp on the 27 mm fixture before pressing **Next**.



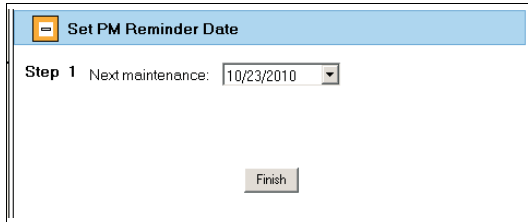
Remove the 27 mm diameter fixture, insert the 127 mm diameter fixture, close the barrel clamp on the fixture. A popup alerts you if the correct size fixture is not installed.





13. Set PM Reminder Date:

The next maintenance reminder date defaults to 12 months. The reminder can be set to an earlier date but it cannot be set to a date past the 12-month default.



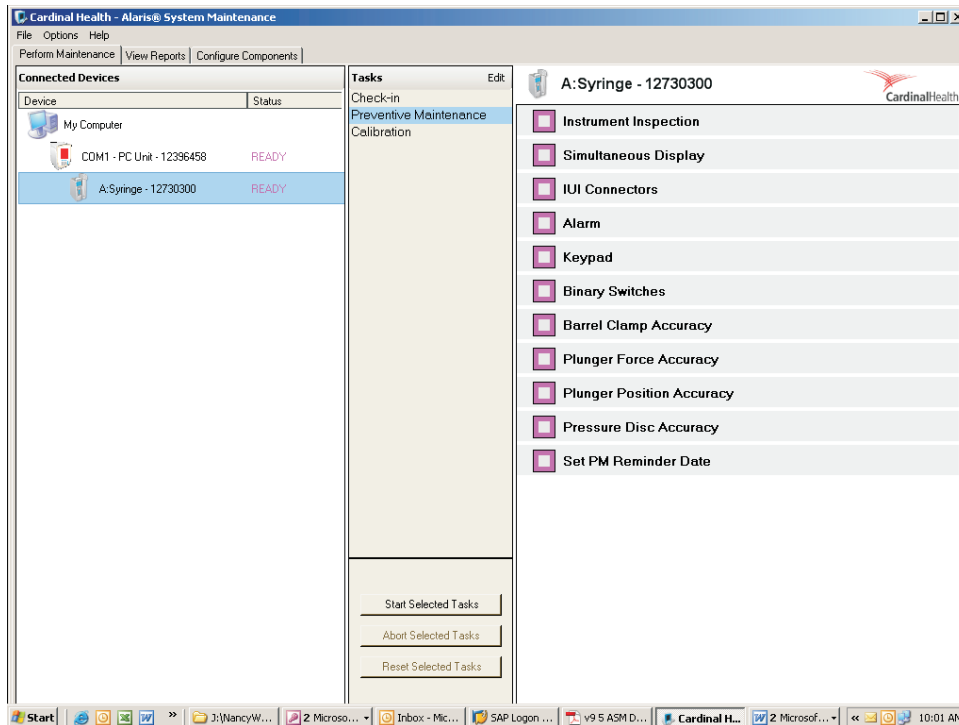
Syringe Module

Test Equipment

Test Equipment	Manufacturer	Model/Part Number	Application
extension set, with pressure disc, small bore	CareFusion	10014917	pressure test
gauge, pressure, digital (peak hold)	Either of the following: <ul style="list-style-type: none"> Heise (www.heise.com) Ashcroft (www.ashcroft.com) 	<ul style="list-style-type: none"> PTE-1 (accuracy from 0.1 to 0.025% span) 2089, 2086, or 2084 (accuracy from $\pm 0.05\%$, 0.10%, or 0.25% of span) or an equivalent gauge with: <ul style="list-style-type: none"> (a) unit of measure in mmHg (b) accuracy of $\pm 1\%$ (c) range of 0-1500 mmHg 	pressure test / calibration
gauge, syringe height	CareFusion	148181-100	plunger position calibration / verification
kit, calibration tools ^a	CareFusion	10010692	all tools required for calibration and verification
kit, force sensor	CareFusion	10010691	force sensor calibration / verification
kit, syringe sizer and height gauge	CareFusion	148182-100	plunger position / barrel size calibration / verification
kit, syringe sizer gauge	CareFusion	148180-100	barrel size calibration / verification
tubing, silicone	CareFusion	303109 or equivalent	pressure test
valve, 3-way	CareFusion	97555 or equivalent	pressure test

- a. To check the spring cartridge verification, preload the cartridge with 0.50 \pm 0.05 lbs. Measure the deflection from the preload point after adding 19.5 lbs additional load (20 \pm 0.05 lbs total). Acceptable deflection is within 0.891-0.940 inches. Use of Instron or equivalent is recommended.

1. In the Connected Devices pane, click the Syringe Module to be tested.
2. Click **Preventive Maintenance** in the Tasks list (double-click to begin testing immediately).



3. Follow the instructions displayed on the screen for each test.

NOTE

See *Syringe Module and PCA Module Tasks* on page 178 for an explanation of each task.

4. Instrument Inspection:

Step 1 Clean any surfaces where solution or obstructions have accumulated.

Examine the instrument for overall condition.

Verify that

1. The case is clean and free from residue, especially near moving parts.
2. Keys, labels, and markings are free of damage and are clearly legible.
3. No tape or other foreign material is on the side of the module.
4. Visually inspect all IUI connectors for damage, bent pins, and signs of corrosion (green material). Replace if found.
5. Case has no apparent damage.
6. IUI latch moves freely.
7. Each key is illuminated by backlight (you may need to cover the keys in order to see the backlight).

Step 2 Examine the instrument for overall condition.

Verify that:

1. The knob opens and closes freely and the claws open and close with it.
2. The plunger head moves up and down freely when the knob is open.
3. The knob, claws, plunger detect, and force sensors have no visible damage.
4. The barrel clamp has no visible damage and moves/rotates freely.
5. The pressure sensor has no visible damage.
6. No other external damage is apparent.
7. The flange and flange sensor have no visible damage.

Step 3 Determine Pass/Fail

Press the 'Pass' button if the test has successfully completed. If any errors have occurred, press the 'Fail' button.

If the test failed, click **Abort Selected Tasks** to discontinue testing. Correct the failure(s) before continuing with the tests.

5. Simultaneous Display:

If you want the test to run continuously, select **Run test in continuous mode**.

Simultaneous Display

Step 1 Run test in continuous mode

Verify:

- 1) Channel ID display.
- 2) Scrolling display.
- 3) Rate display
- 4) Lighthouse LED's
- 5) No missing pixels

Note that the test may take a few moments to complete after clicking 'Stop'. If running the test in continuous mode, uncheck the continuous mode checkbox or press 'Stop' to stop the test.

When ready to continue, press the 'Next' button.

- **Channel ID display**—all segments must light.
- **Scrolling display** and **Rate display**—must flash dashes and zeros.
- **Lighthouse LED's**—all LEDs must light.

Step 2 Determine Pass/Fail

Press the 'Pass' button if the test has successfully completed. If any errors have occurred, press the 'Fail' button.

6. IUI Connectors:

IUI Connectors

Step 1 Visually inspect all IUI connectors for damage, bent pins, and signs of corrosion (green material). Replace if found.

Connect one other module (not an Auto-ID module) to the Syringe, 'sandwiching' the Syringe (if not already done). Remove any other modules other than the Syringe, the PC Unit, and the module on the other side of the Syringe

Wait for the Channel ID to appear on the attached units before pressing 'Next'.

'Sandwiching'—the module being tested is connected between the PC Unit and another module.

Step 2 Result of device poll

Summary of the number of devices detected

Expected number of devices :	3
Actual Number of devices :	3
	Pass

Press 'Finish' to continue

7. Alarm:

Alarm

Step 1 Do you see the Syringe Alarm LED and hear the PC Unit speaker alarming?

If the audio is weak, check the audio volume setting (refer to applicable Alaris® System DFU).

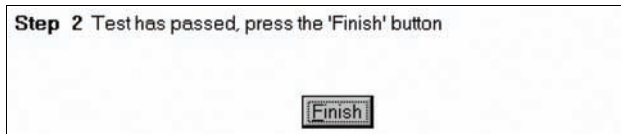
Step 2 A Pass result will be entered for this device.

8. Keypad:



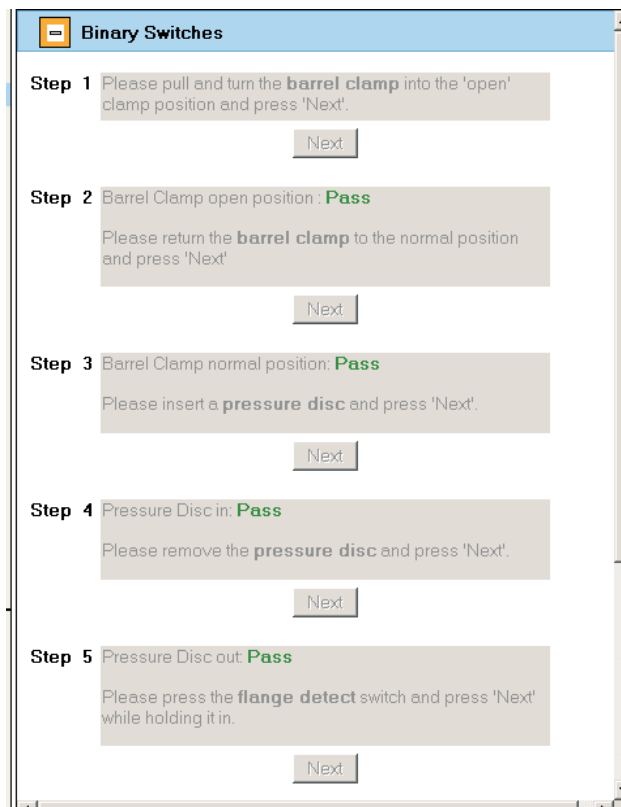
Press each key one time.

- Green text—the key was pressed one time and passed the test.
- Red text—the key was pressed two times, or it was pressed one time and failed the test. If the text is red because you pressed the key more than one time, you can indicate that the test has passed.



9. Binary Switches:

The Binary Switches test verifies that the buttons and switches on the Syringe Module are open/closed, on/off, pushed/not pushed, and so on.



Step 6 Flange Detect in: **Pass**

Please release the **flange detect** switch and press 'Next'.

Next

Step 7 Flange Detect out: **Pass**

Please open the claws with the **knob** and press 'Next' while holding them open.

Next

Step 8 Knob open: **Pass**
Split Nut open: **Pass**

Please release the **knob** allowing the claws to close and press 'Next'.

Next

Step 9 Knob closed: **Pass**
Split Nut closed: **Pass**

Please press the **plunger detect** switch and press 'Next' while holding it in.

Next

Step 10 Plunger Detect in: **Pass**

Please release the **plunger detect** switch and press 'Next'.

Next

Step 11

Pressure Disc	Present: Pass	Absent: Pass
Plunger	Pressed: Pass	Released: Pass
Flange	Pressed: Pass	Released: Pass
Knob	Open: Pass	Closed: Pass
Split Nut	Open: Pass	Closed: Pass
Barrel Clamp	Open: Pass	Normal: Pass

Switch Sensor Test Results **Pass**

Finish

10. Barrel Clamp Accuracy:

Barrel Clamp Accuracy

Step 1 Please insert and clamp the 15 mm diameter fixture.

When ready, press the 'Next' button to begin the verification.

Next

Step 2 Please insert and clamp the 25 mm diameter fixture.

When ready, press the 'Next' button to continue the verification.

Step 3 Please remove the fixture

Barrel Size Ver. Results	Passed
15 mm Verification:	Passed
25 mm Verification:	Passed

The device is functioning properly.

11. Plunger Force Accuracy:

Plunger Force Accuracy

Step 1 Please insert the spring calibration fixture into the unit.

Close the barrel clamp, but do not attach the plunger header to the calibration fixture.

When ready, press the 'Next' button to take initial readings.

IMPORTANT: To avoid damage to the plunger header, ensure that the plunger header is not attached to the calibration fixture.

Step 2 Please attach the plunger head to the calibration fixture.


To prevent damage to the unit, do not move the plunger head while the test is in progress until instructed to do so.

When ready, press the 'Next' button to begin the verification.

IMPORTANT: To avoid instrument damage, do not move the plunger head during this test.

Step 3 Verification Progress

Adjusting to starting position



Step 4 When ready, press the 'Next' button to continue the verification.

Step 5 Release all force from the system and remove the gauge

Force Verification Results	Passed
11.6 lb Verification:	Passed
3.0 lb Verification:	Passed

The device is functioning properly.

12. Plunger Position Accuracy:

Plunger Position Accuracy

Step 1 Please insert the 27 mm fixture, adjusting the plunger head to touch the top of the fixture, ensuring that the plunger detect switch is fully depressed.

Do not use a pressure disc for this test.

When ready, press the 'Next' button to begin the verification.

Step 2 Taking Measurements

Adjusting plunger position backwards

Step 3 Please insert the 127 mm fixture, adjusting the plunger head to touch the top of the fixture, ensuring that the plunger detect switch is fully depressed.

When ready, press the 'Next' button to continue the verification.

Step 4 Taking Measurements

Adjusting plunger position backwards

Step 5 Please remove the fixture.

Plunger Position Ver. Results **Passed**

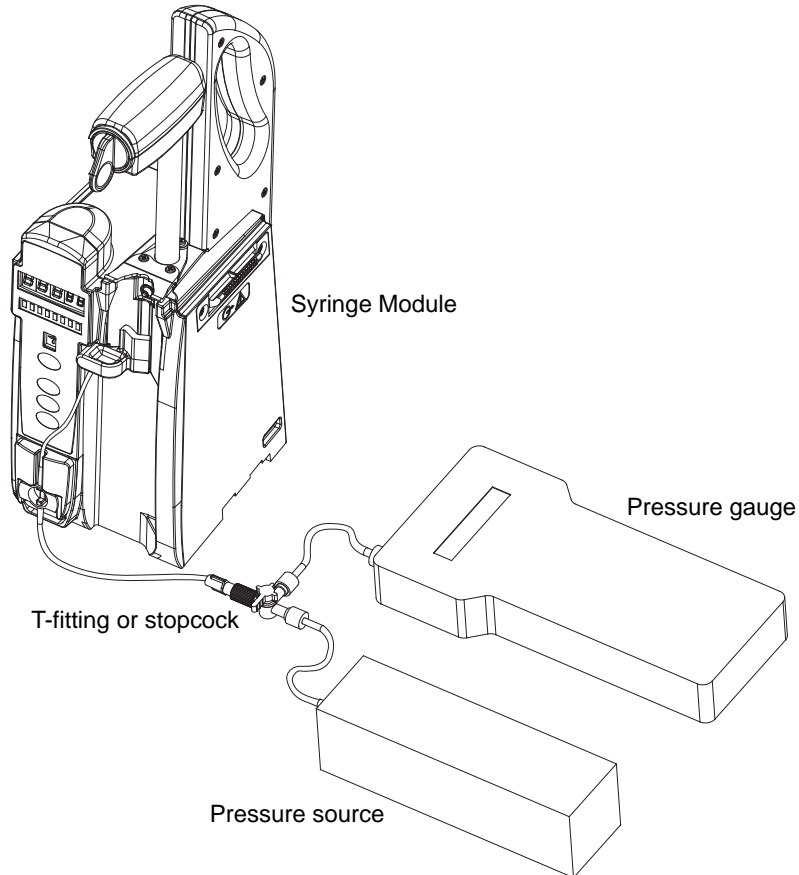
27 mm Calibration: **Passed**

127 mm Calibration: **Passed**

The device is functioning properly.

13. Pressure Disc Accuracy:

Set up the Syringe Module for the pressure test.

Pressure Test/Calibration Setup

Pressure Disc Accuracy

Step 1 Verify that the pressure gauge is set to mmHg. Install the test set in to the module and connect the set to the pressure gauge.

Apply pressure until the pressure gauge reads **250 mmHg** +/- 5 mmHg.

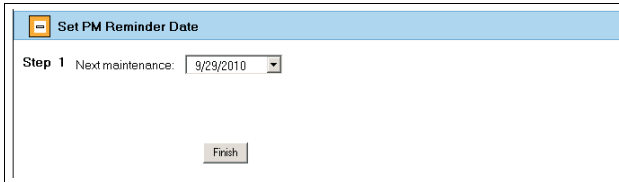
Step 2 Pressure Verification Results **Passed**

250 mmHg Verification: **Passed**

The device is functioning properly.

14. Set PM Reminder Date:

The next maintenance reminder date defaults to 12 months. The reminder can be set to an earlier date but it cannot be set to a date past the 12-month default.



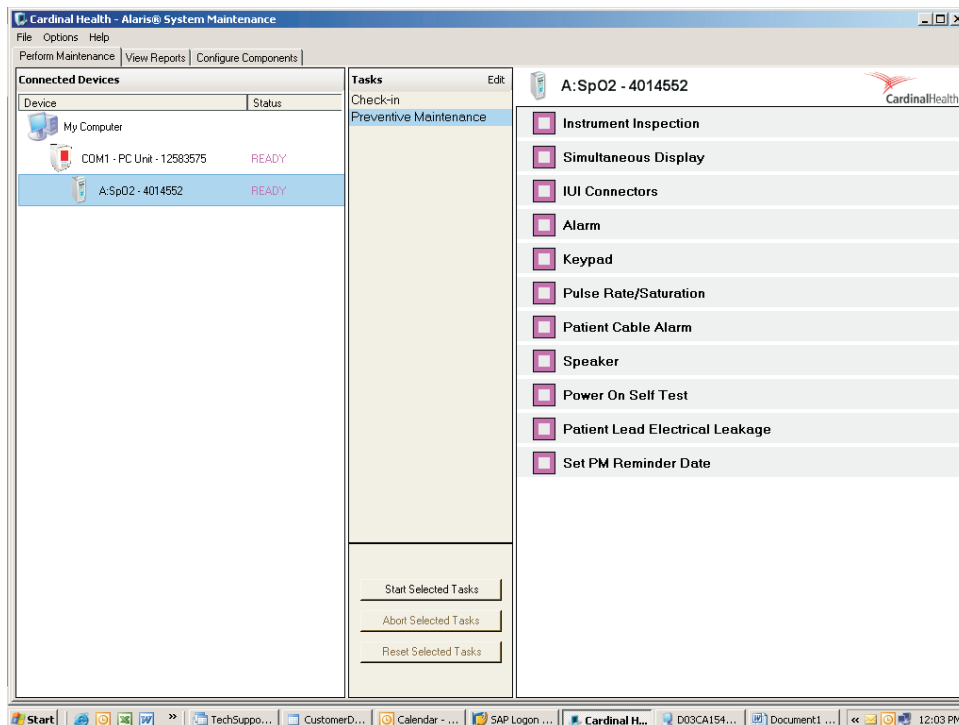
The screenshot shows a dialog box titled "Set PM Reminder Date". It contains a single step labeled "Step 1" with the text "Next maintenance:" followed by a dropdown menu showing the date "9/29/2010". At the bottom center of the dialog box is a button labeled "Finish".

SpO₂ Module

Test Equipment

Test Equipment	Manufacturer	Model Number	Application
Analyzer, electrical	Fluke BioMedical	232D or equivalent	patient lead electrical leakage
Cable, Masimo® Simulator	Clinical Dynamics	SS-MAS	pulse rate/saturation
Cable, Nellcor® Oximax® Simulator	Clinical Dynamics	SS-NEL	pulse rate/saturation
SpO ₂ simulator	Clinical Dynamics	SmartSat Pulse Oximetry Analyzer SS-100A, or equivalent	pulse rate/saturation

1. In the Connected Devices pane, click the SpO₂ Module to be tested.
2. Click **Preventive Maintenance** in the Tasks list (double-click to begin testing immediately).



3. Follow the instructions displayed on the screen for each test.

NOTE

See *SpO₂ Module Tasks* on page 182 for an explanation of each task.

4. Instrument Inspection:

Step 1 Clean any surfaces where solution or obstructions have accumulated.

Examine the instrument for overall condition.

Verify that:

1. The case is clean and free from residue, especially near moving parts.
2. Keys, labels, and markings are free of damage and are clearly legible.
3. No tape or other foreign material is on the side of the module.
4. Visually inspect all IUI connectors for damage, bent pins, and signs of corrosion (green material). Replace if found.
5. Case has no apparent damage.
6. IUI latch moves freely.
7. Each key is illuminated by backlight (you may need to cover the keys in order to see the backlight).

Step 2 Examine the instrument for overall condition.

Verify that:

1. The SpO₂ sensor port is in good condition and there are no obstructions.
2. No other external damage is apparent.

Step 3 Determine Pass/Fail

Press the 'Pass' button if the test has successfully completed. If any errors have occurred, press the 'Fail' button.

If the test failed, click **Abort Selected Tasks** to discontinue testing. Correct the failure(s) before continuing with the tests.

5. Simultaneous Display:

If you want the test to run continuously, click the **Run test in continuous mode** checkbox.

Step 1 Run test in continuous mode

Verify:

- 1) Channel ID display.
- 2) Scrolling display.
- 3) Pulse display
- 4) Saturation display
- 5) Pleth bar graph display
- 6) Lighthouse LED's
- 7) No missing pixels

Note that the test may take a few moments to complete after clicking 'Stop'. If running the test in continuous mode, uncheck the continuous mode checkbox or press 'Stop' to stop the test.

When ready to continue, press the 'Next' button.

- **Channel ID display**—all segments must light.
- **Scrolling display, Pulse display, and Saturation display**—must flash dashes and zeros.
- **Pleth bar graph display**—all segments must light.
- **Lighthouse LED's**—all LEDs must light.

Step 2 Determine Pass/Fail
Press the 'Pass' button if the test has successfully completed. If any errors have occurred, press the 'Fail' button.

Pass Fail

6. IUI Connectors:

IUI Connectors

Step 1 Visually inspect all IUI connectors for damage, bent pins, and signs of corrosion (green material). Replace if found.

Connect one other module (not an Auto-ID module) to the SpO₂, 'sandwiching' the SpO₂ (if not already done). Remove any other modules other than the SpO₂, the PC Unit, and the module on the other side of the SpO₂

Wait for the Channel ID to appear on the attached units before pressing 'Next'.

Next

'Sandwiching'—the module being tested is connected between the PC Unit and another module.

Step 2 Result of device poll
Summary of the number of devices detected

Expected number of devices :	3
Actual Number of devices :	3
	Pass

Press 'Finish' to continue

Finish

7. Alarm

Alarm

Step 1 Do you see the SpO₂ Alarm LED and hear the PC Unit speaker alarming?

Yes No

If the audio is weak, check the audio volume setting (refer to applicable Alaris® System DFU).

Step 2 A Pass result will be entered for this device.

Finish

8. Keypad:

Keypad

Step 1 Please press every key and verify the key pressed.

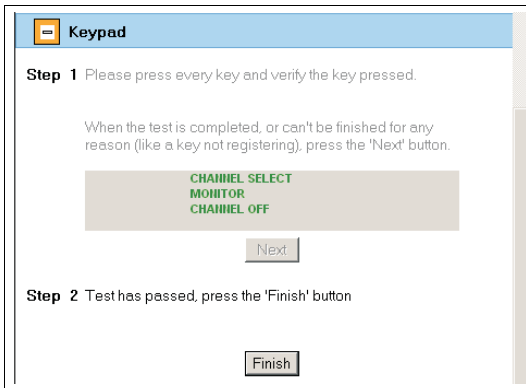
When the test is completed, or can't be finished for any reason (like a key not registering), press the 'Next' button.

CHANNEL SELECT
MONITOR
CHANNEL OFF

Next

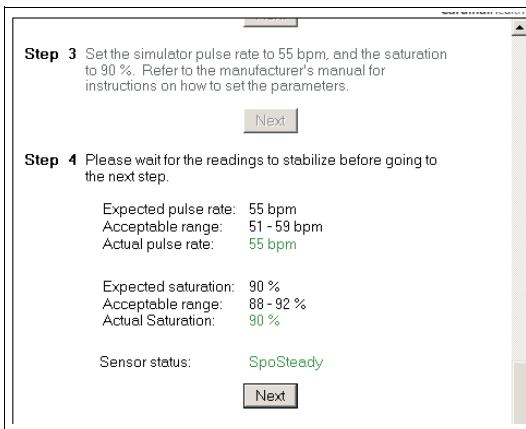
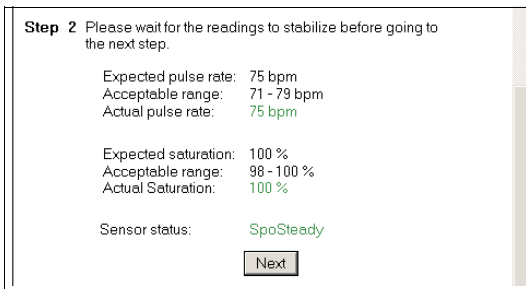
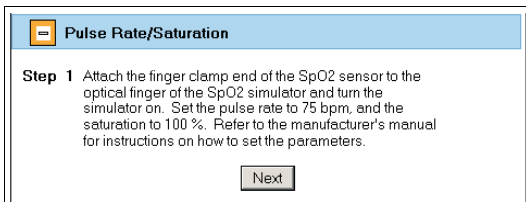
Press each key one time.

- Green text—the key was pressed one time and passed the test.
- Red text—the key was pressed two times, or it was pressed one time and failed the test. If the text is red because you pressed the key more than one time, you can indicate that the test has passed.



9. Pulse Rate/Saturation:

If the Clinical Dynamics SmartSat is being used, with a direct feed into the device, connect it before starting the test.



Step 5 The overall **PASSED** status and the following details will be entered for this device.

Reading 1	
Pulse	-PASSED
Saturation	-PASSED
Reading 2	
Pulse	-PASSED
Saturation	-PASSED

Finish

10. Patient Cable Alarm:

If the Clinical Dynamics SmartSat is being used, use a finger sensor instead of the patient cable.

Patient Cable Alarm

Step 1 Connect the patient cable to the SpO₂.

Next

Step 2 Waiting for the SpO₂ to Initialize...

Step 3 Disconnect the patient cable.

Next

Step 4 Re-connect the patient cable.

Next

Step 5

Initialization	Passed
Disconnect	Passed
Connect	Passed

The preceding information will be entered into the database.

Finish

11. Speaker:

Speaker

Step 1 Is the SpO₂ speaker sounding?

Yes **No**

Step 2 A Pass result will be entered for this device.

Finish

12. Power On Self Test:

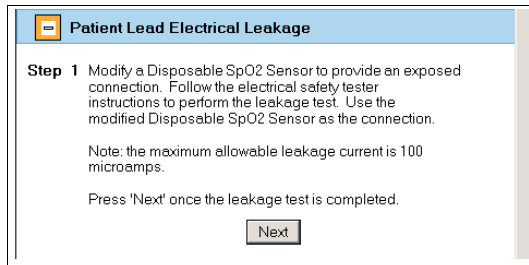
Power On Self Test

Step 1 The POST test has passed. A Pass result will be entered for this device.

Finish

13. Patient Lead Electrical Leakage:

The System Maintenance Software is not used to perform this test. The test can be performed before beginning Preventive Maintenance testing or at this point in the testing.

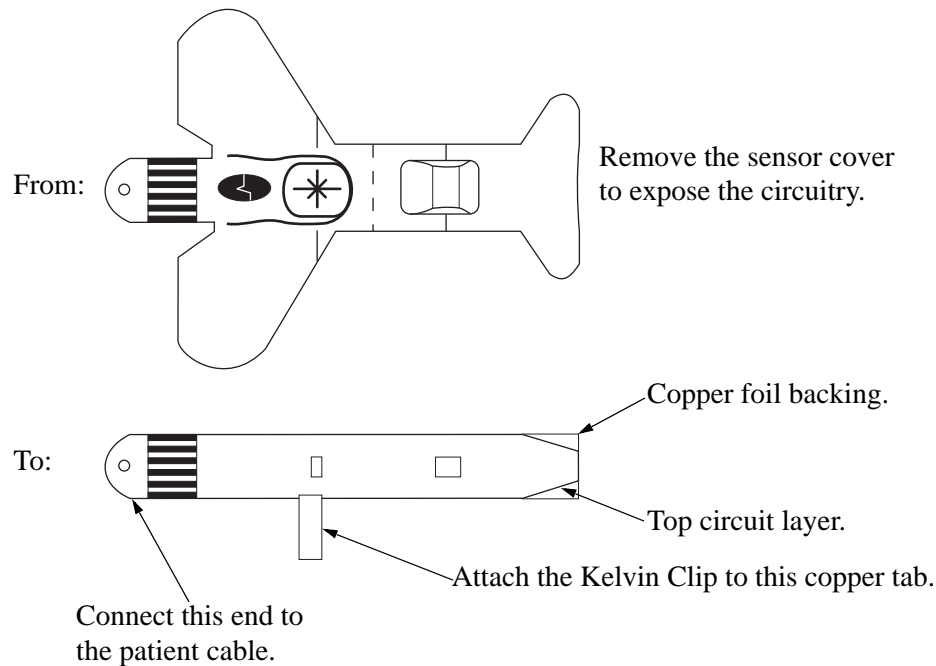


Use a Fluke Biomedical Model 232D or equivalent electrical safety analyzer to measure the electrical leakage current. Refer to the enclosure or case leakage instructions in the test equipment's operation manual for the proper measurement technique.

Use the following illustration to modify and use a disposable SpO₂ Sensor for the test. The maximum allowable leakage current is 100 μ A, for normal and reversed line polarity.

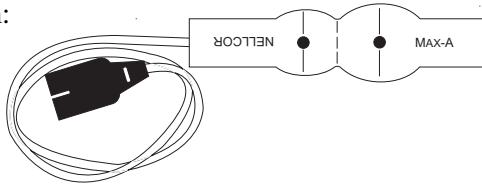
Disposable SpO₂ Sensor Modification and Test Point

Masimo® Sensor



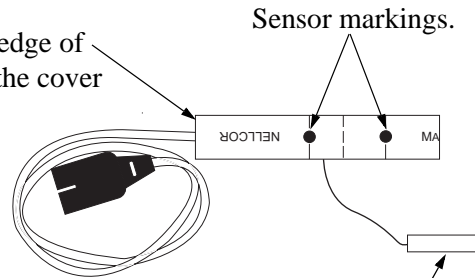
Nellcor® Sensor

From:



To:

Carefully cut around the edge of the sensor and pull back the cover to expose the circuitry.



Pull the copper tab loose. Reseal the cover. Ensure that the sensors are lined up between the sensor windows and markings.

Attach the Kelvin Clip to this copper tab.

Step 2 Select the result to be entered into the database.

1. Set PM Reminder Date:

The next maintenance reminder date defaults to 12 months. The reminder can be set to an earlier date but it cannot be set to a date past the 12-month default.

Set PM Reminder Date

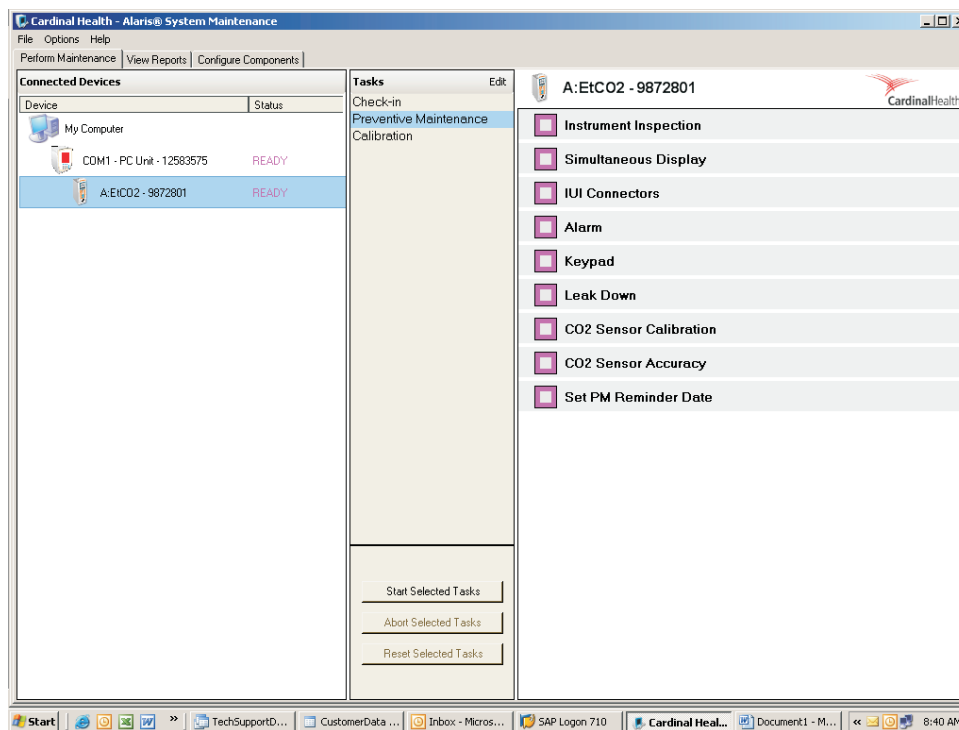
Step 1 Next maintenance: 10/23/2010

EtCO₂ Module

Test Equipment

Test Equipment	Manufacturer	Model Number	Application
EtCO ₂ pod (disposable)	Microstream®	007768	flow test/calibration
flow meter	Dwyer (www.dwyer-inst.com.au)	VFA-22, or equivalent	flow test/calibration
gas	Scott Medical	0304653SROBD, or equivalent gas with CO ₂ content of 4-6%	flow test/calibration
pressure transducer simulator/tester	Utah Medical	Delta-Cal 650-95,0 or equivalent	leak-down test
tubing, silicone	CareFusion	303109 or equivalent	pressure test
valve, 3-way	CareFusion	97555 or equivalent	pressure test

1. In the Connected Devices pane, click the EtCO₂ Module to be tested.
2. Click **Preventive Maintenance** in the Tasks list (double-click to begin testing immediately).



3. Follow the instructions displayed on the screen for each test.

NOTE

See *EtCO₂ Module Tasks* on page 183 for an explanation of each task.

4. Instrument Inspection:

Step 1 Clean any surfaces where solution or obstructions have accumulated.

Examine the instrument for overall condition.

Verify that:

1. The case is clean and free from residue, especially near moving parts.
2. Keys, labels, and markings are free of damage and are clearly legible.
3. No tape or other foreign material is on the side of the module.
4. Visually inspect all IUI connectors for damage, bent pins, and signs of corrosion (green material). Replace if found.
5. Case has no apparent damage.
6. IUI latch moves freely.
7. Each key is illuminated by backlight (you may need to cover the keys in order to see the backlight).

Step 2 Examine the instrument for overall condition.

Verify that:

1. The rotary door moves freely.
2. When opened and released, the rotary door closes freely.
3. There is no visible obstruction or blockage where the disposable attaches.
4. No other external damage is apparent.

Step 3 Determine Pass/Fail

Press the 'Pass' button if the test has successfully completed. If any errors have occurred, press the 'Fail' button.

If the test failed, click **Abort Selected Tasks** to discontinue testing. Correct the failure(s) before continuing with the tests.

5. Simultaneous Display:

If you want the test to run continuously, click the **Run test in continuous mode** checkbox.

Step 1 Run test in continuous mode

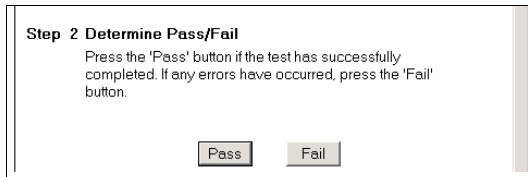
Verify:

- 1) Channel ID display.
- 2) Scrolling display.
- 3) ETCO₂ display
- 4) Respiration rate display
- 5) Lighthouse LED's
- 6) No missing pixels

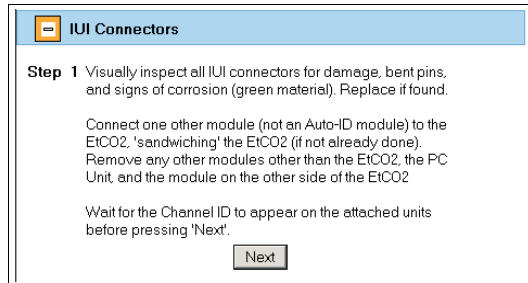
Note that the test may take a few moments to complete after clicking 'Stop'. If running the test in continuous mode, uncheck the continuous mode checkbox or press 'Stop' to stop the test.

When ready to continue, press the 'Next' button.

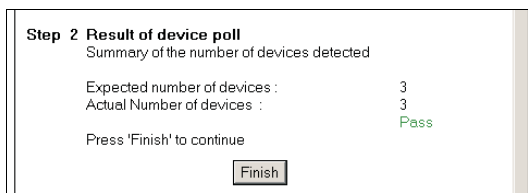
- **Channel ID display**—all segments must light.
- **Scrolling display, ETCO₂ display, and Respiration rate display**—must flash dashes and zeros.
- **Lighthouse LED's**—all LEDs must light.



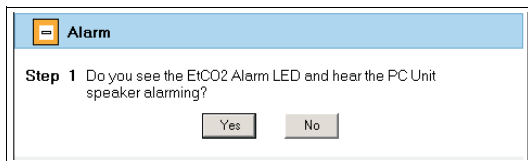
6. IUI Connectors:



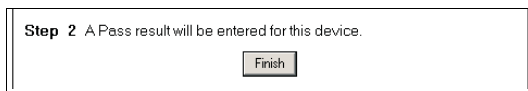
'Sandwiching'—the module being tested is connected between the PC Unit and another module.



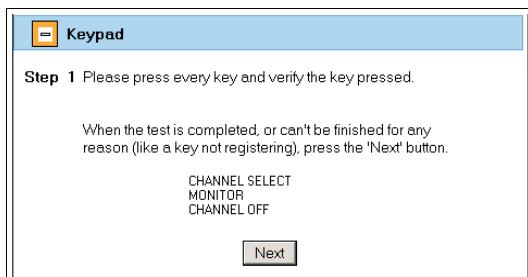
7. Alarm:



If the audio is weak, check the audio volume setting (refer to applicable Alaris® System DFU).

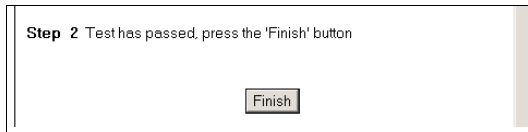


8. Keypad:



Press each key one time.

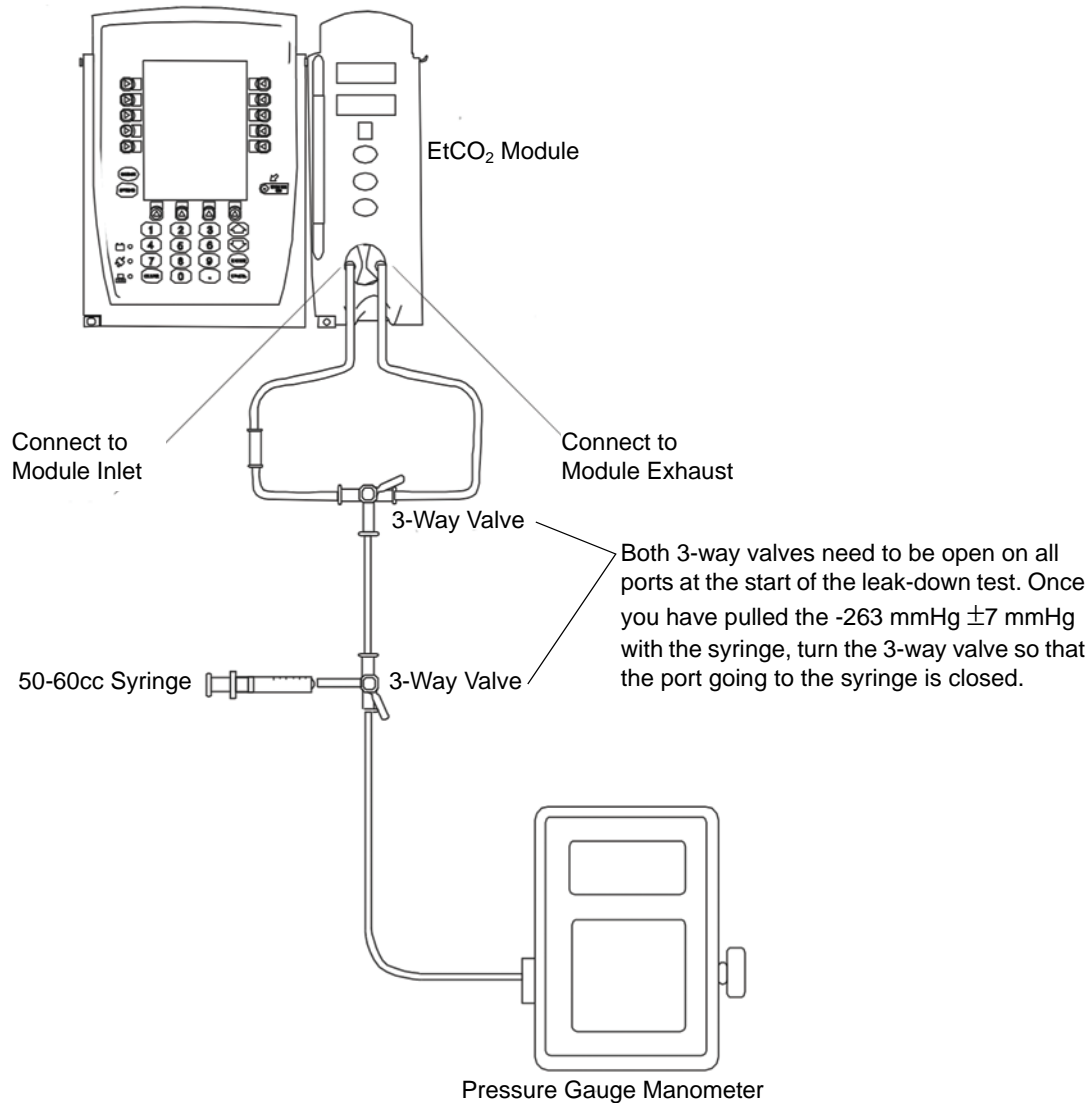
- Green text—the key was pressed one time and passed the test.
- Red text—the key was pressed two times, or it was pressed one time and failed the test. If the text is red because you pressed the key more than one time, you can indicate that the test has passed.



9. Leak Down:

Leak-Down Test

Ensure that the vacuum gauge is set to mmHg. Create a test harness to match the components that are used in the facility.




Leak Down

Step 1 Please connect the leak down test harness to the EtCO₂ Module, vacuum source and pressure/vacuum gauge. Ensure the vacuum gauge is set to read mmHg and zero the gauge. Apply a vacuum of -263 mmHg +/-7 mmHg. Record the actual applied vacuum, seal the system if appropriate, and when ready, press the 'Next' button.


If you get a vacuum leak, pinch off the tubing at the inlet and at the exhaust, to ensure that the leak is not in the setup.

Step 2 Waiting for 10 seconds: 10



Step 3 After pressing the 'Next' button the system will stabilize. When the timer completes, record the vacuum remaining in the system. Subtract this value from the initial to determine vacuum lost. When ready to continue, press the 'Next' button.

Step 4 Waiting for 60 seconds: 60



Step 5 Enter the vacuum lost from the system. With the test complete, please disconnect the disposable before pressing 'Next'.

Vacuum change: mmHg

Passing values are 0 to 30 mmHg

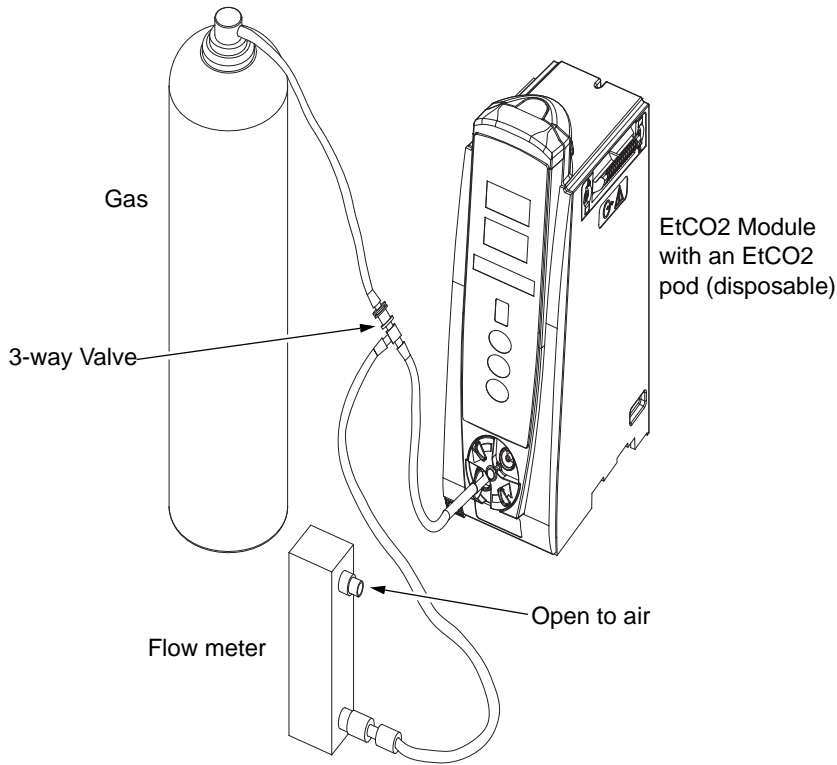
The **Vacuum change** entry is the value calculated in step 3 and rounded to a whole number.

Step 6 Test Result: Passed

10. CO₂ Sensor Calibration:

Set up the EtCO₂ Module for the flow test/calibration.

Flow Test/Calibration Setup



CO₂ Sensor Calibration

Step 1 Reminder: Do not calibrate this unit unless it has been warming up for at least 20 minutes.

Please connect a disposable and attach the gas source. When instructed, start the gas flowing and leave it flowing until instructed to stop.

Caution: The gas flow rate must be above 500 mL/min. Use of a flow meter is recommended.

Next

500 mL/min is equivalent to 0.5 liters per minutes. Vent the flow meter to open air.

Step 2 Enter the percentage of CO₂ in the calibration gas (there should be 21 % O₂ and the balance should be N₂).

% CO₂

Valid values are 4% to 6%

Next

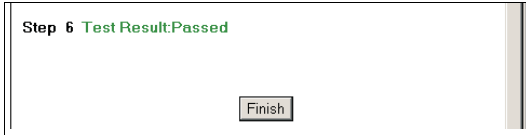
Step 3 Start the gas flow

Next

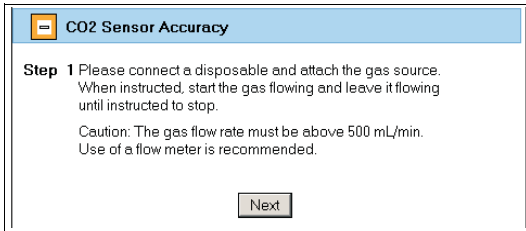
Do not turn off the gas flow until instructed to in step 5.

Step 4 Running calibration. Keep the gas flowing.

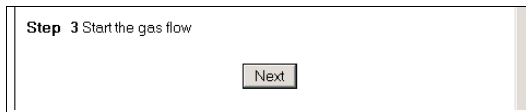
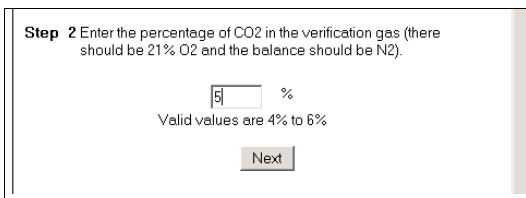
Step 5 Calculating... Please turn off the gas source



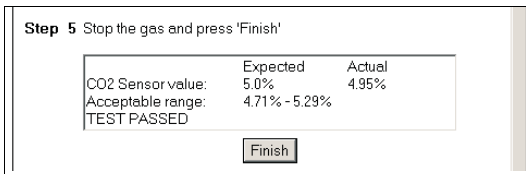
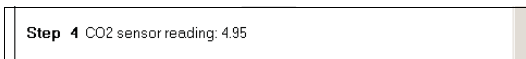
11. CO2 Sensor Accuracy:



500 mL/min is equivalent to 0.5 liters per minutes. Vent the flow meter to open air.

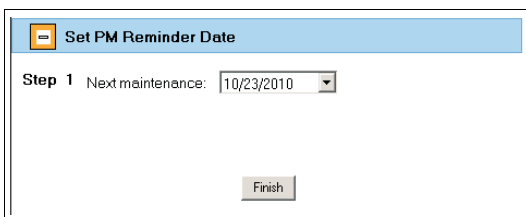


Do not turn off the gas flow until instructed to in step 5.



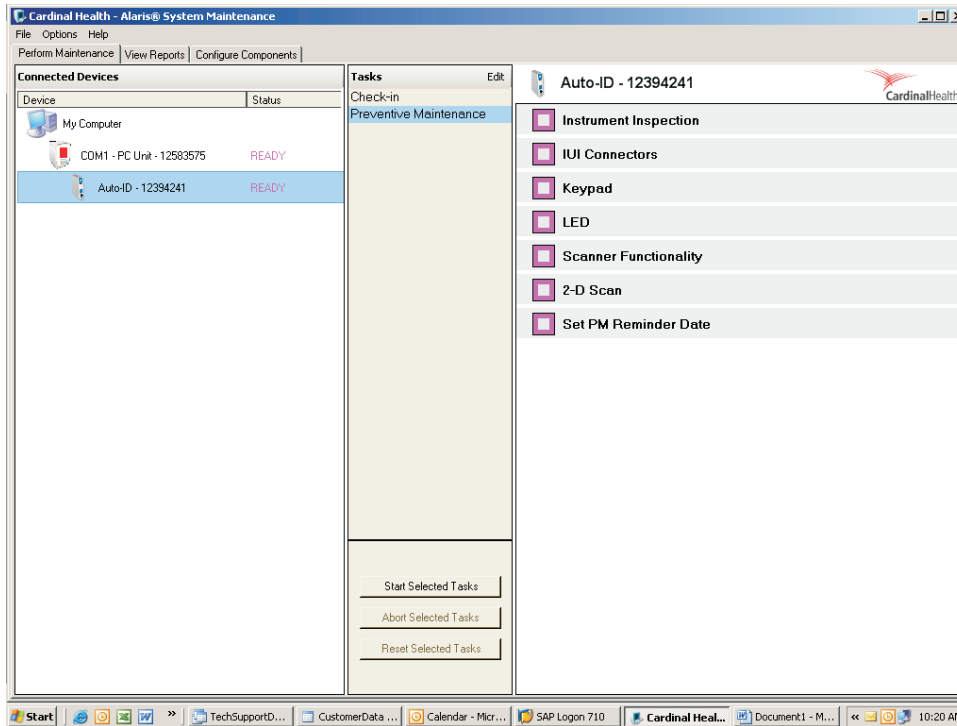
12. Set PM Reminder Date:

The next maintenance reminder date defaults to 12 months. The reminder can be set to an earlier date but it cannot be set to a date past the 12-month default.



Auto-ID Module and Handheld Scanner

1. In the Connected Devices pane, click the Auto-ID Module to be tested.
2. Click **Preventive Maintenance** in the Tasks list (double-click to begin testing immediately).

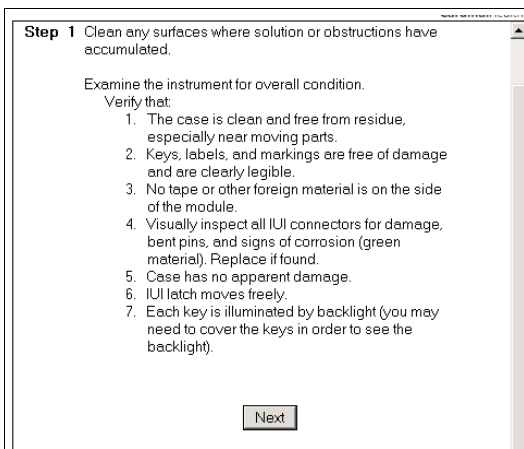


3. Follow the instructions displayed on the screen for each test.

NOTE

See *Auto-ID Module and Handheld Scanner Tasks* on page 184 for an explanation of each task.

4. Instrument Inspection:



Step 2 Examine the instrument for overall condition.

Verify that:

1. The lense is in good condition (free from scratches and cracks).
2. The lense is free from obstructions, such as tape or paint.
3. No other external damage is apparent.

Step 3 Determine Pass/Fail

Press the 'Pass' button if the test has successfully completed. If any errors have occurred, press the 'Fail' button.

If the test failed, click **Abort Selected Tasks** to discontinue testing. Correct the failure(s) before continuing with the tests.

5. IUI Connectors:

IUI Connectors

Step 1 Visually inspect all IUI connectors for damage, bent pins, and signs of corrosion (green material). Replace if found.

Connect one other module (not an Auto-ID module) to the Auto-ID, 'sandwiching' the Auto-ID (if not already done). Remove any other modules other than the Auto-ID, the PC Unit, and the module on the other side of the Auto-ID.

Wait for the Channel ID to appear on the attached units before pressing 'Next'.

'Sandwiching'—the module being tested is connected between the PC Unit and another module.

Step 2 Result of device poll

Summary of the number of devices detected

Expected number of devices :	2
Actual Number of devices :	2
	Pass

Press 'Finish' to continue

6. Keypad:

Keypad

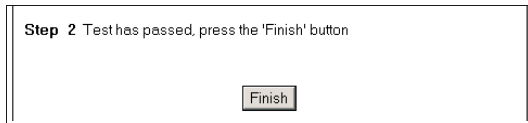
Step 1 Please press every key and verify the key pressed.

When the test is completed, or can't be finished for any reason (like a key not registering), press the 'Next' button.

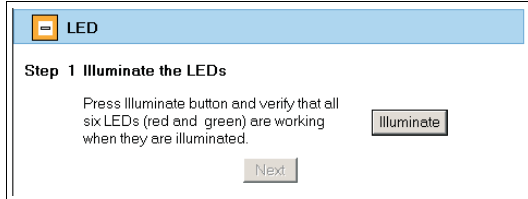
SCAN/CANCEL

Press each key one time.

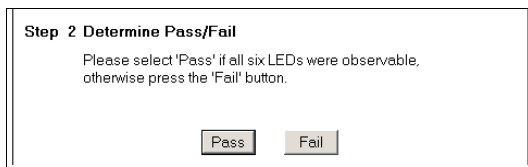
- Green text—the key was pressed one time and passed the test.
- Red text—the key was pressed two times, or it was pressed one time and failed the test. If the text is red because you pressed the key more than one time, you can indicate that the test has passed.



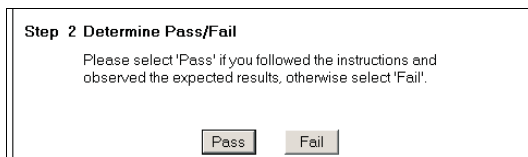
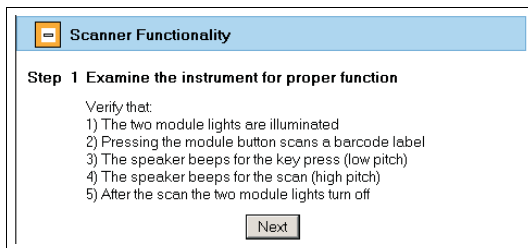
7. LED:



- Red LEDs—Camera lens.
- Green LEDs—Ready.

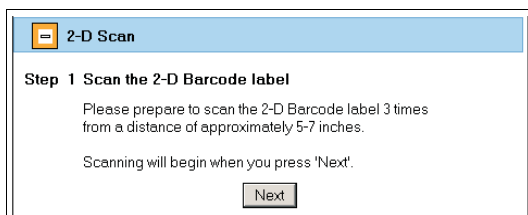


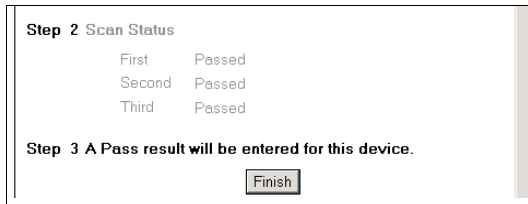
8. Scanner Functionality:



9. 2-D Scan:

See *Auto-ID Module and Handheld Scanner Tasks* on page 184 to locate the 2-D barcode label. For the best results when scanning the barcode, hold the barcode at an approximate 45 degree angle to the scanner.





Step 2 Scan Status

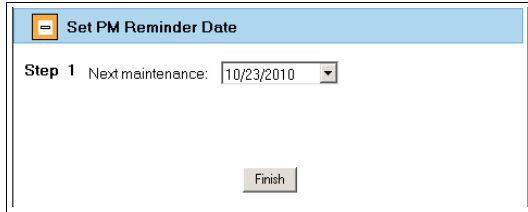
First	Passed
Second	Passed
Third	Passed

Step 3 A Pass result will be entered for this device.

Finish

10. Set PM Reminder Date:

The next maintenance reminder date defaults to 12 months. The reminder can be set to an earlier date but it cannot be set to a date past the 12-month default.



Set PM Reminder Date

Step 1 Next maintenance: 10/23/2010

Finish

Chapter 7

Data Sets and Alerts

Transfer a Data Set

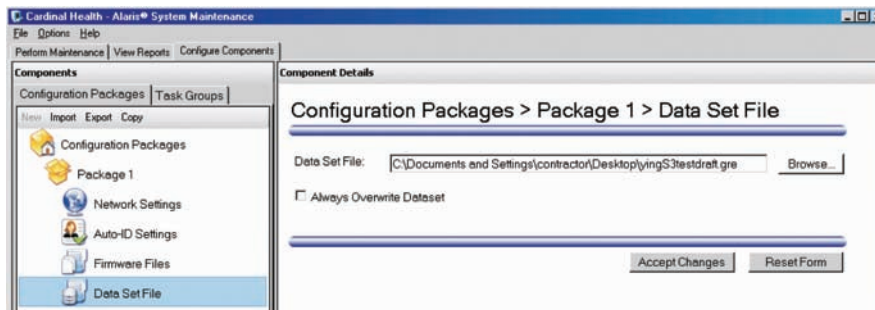
A Data Set can be transferred to a PC Unit through a wireless or serial network.

Use this procedure to transfer a Data Set to a PC Unit. Creating and editing a Data Set is beyond the scope of this manual (see the user manual for the Editor Software).

CAUTION

Users of the Auto-ID Module must download the Data Set wirelessly for the system to properly read and decode bar code labels. Users who do not use the Auto-ID Module can use the following procedure or download the Data Set wirelessly if they are a network user.

1. Associate a Data Set with a configuration package.
 - a. Click the **Configure Components** tab.
 - b. Click the **Configuration Packages** subtab.
 - c. Click **New**.
 - d. Click **Data Set File**.



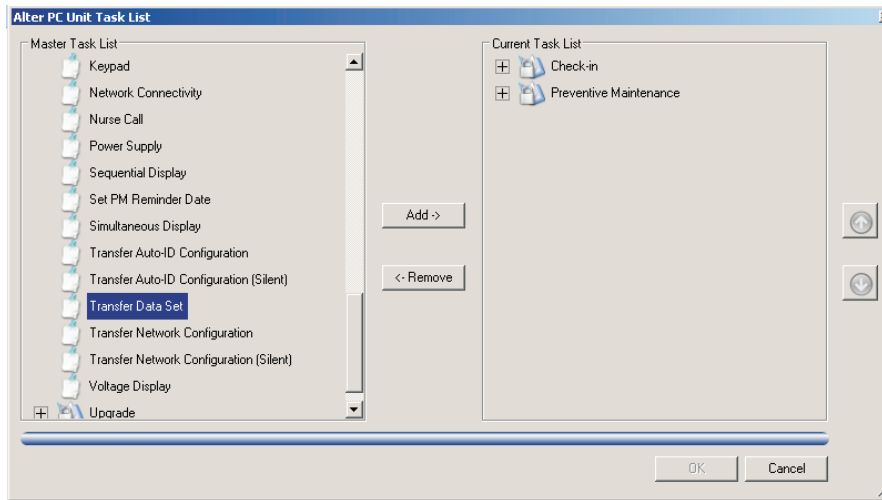
2. Click **Browse** and select an existing Data Set file.
3. Click **Accept Changes**.

This allows the System Maintenance Software to determine which Data Set should be transferred to the PC Unit. Note that the application can maintain multiple packages, each of which is associated with a Data Set. Each Data Set can contain multiple profiles, which are typically used for different types of patients.

At any time, only one Data Set is active on a PC Unit; however, two Data Sets can exist on a PC Unit: the current Data Set and a pending Data Set. A current Data Set is active and used with a current patient. When a new Data Set is transferred to a PC Unit, it becomes a pending Data Set and is not used until you specify that a new patient is being served (by using front panel interface of the PC Unit) and power the PC Unit off and on.

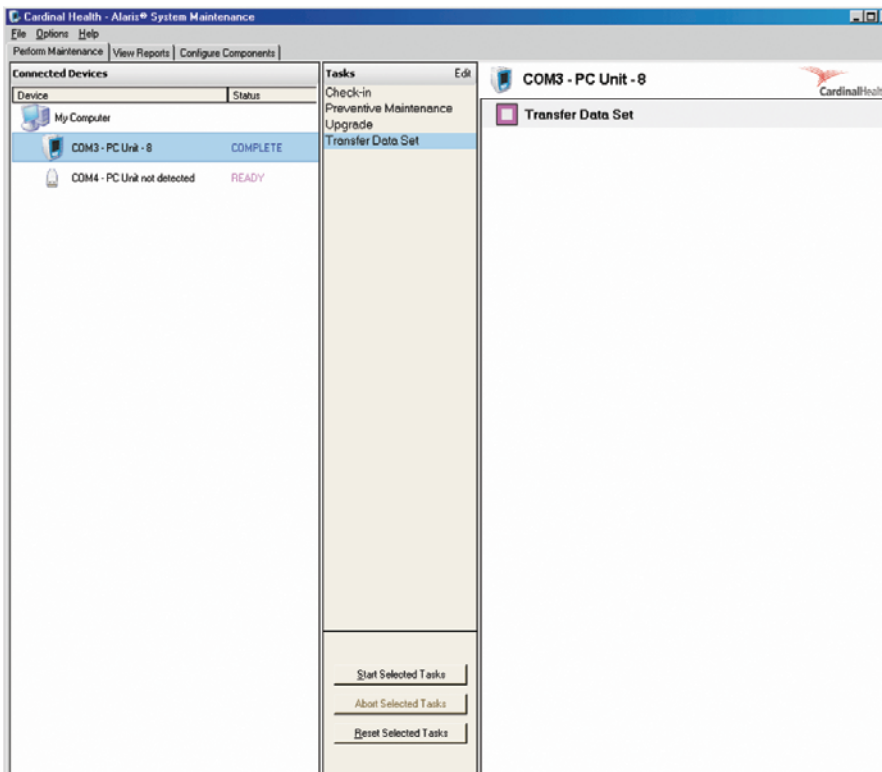
4. To transfer the Data Set, click the **Perform Maintenance** tab and then click the PC Unit in the Connected Devices pane.
5. If **Transfer Data Set** task is not available in the Tasks list, click **Edit** (see *Tasks List* on page 168 for detailed procedures on how to edit the Tasks list).

- In the Alter PCU Task List dialog box, click **Transfer Data Set**.



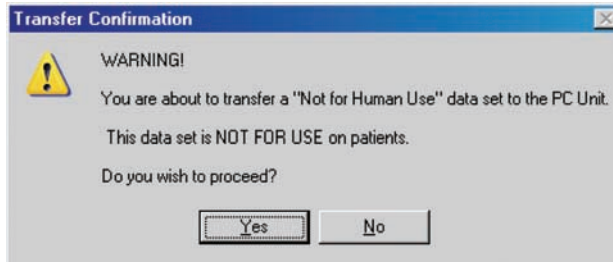
- Click **Add** and then click **OK**.

You can now run the **Transfer Data Set** task for the PC Unit and transfer the Data Set specified earlier in the **Configuration Packages** tab.



8. In the Tasks list, click **Transfer Data Set** and then click **Start Selected Tasks**.

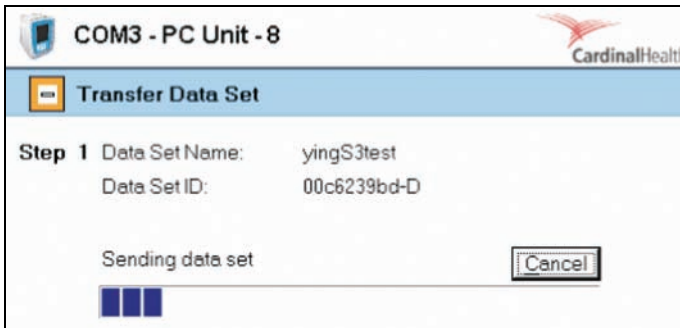
If you are using a Data Set that has not been released, the following warning message is displayed.



NOTE

If the Data Set is not compatible with the embedded software, an error message appears.

As the Data Set is transferred to the PC Unit, a progress bar indicates the transfer status. Transfer usually only takes a few seconds.

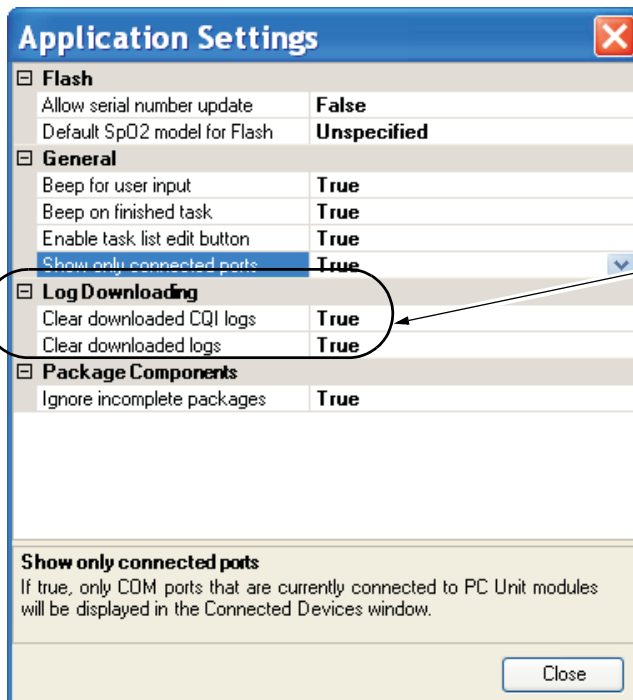


Download Alerts

Alerts are stored in the CQI log on the PC Unit. Typically, you need to periodically download these alerts to the CQI database. The general topic of CQI alerts is beyond the scope of this manual. Refer to the software user manual for CQI Reporter.

You should first determine if you wish to manually or automatically clear the CQI log stored on the PC Unit after downloading the alerts from it. The PC Unit has a limit on the number of CQI alerts it can store. Therefore, it is important to clear the CQI log after you download alerts from it, unless you are downloading for testing purposes and want to retain the alerts on the device.

1. On the main menu, click **Options > Application Settings**.

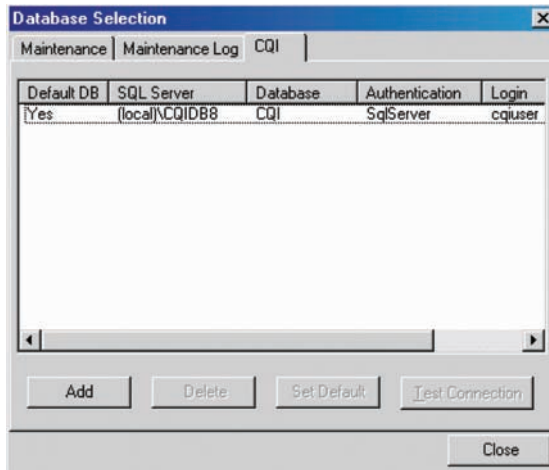


The default setting to clear downloaded logs is **True**. This setting *automatically clears* the logs during the log download.

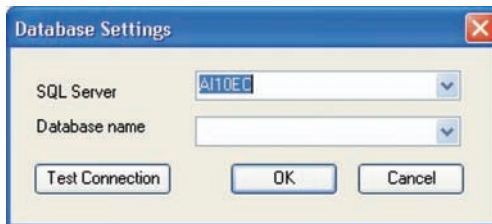
To allow the logs to be *manually cleared* after downloading, change the setting to **False**.

2. Click **True** to automatically clear logs during download or **False** if logs are to be manually cleared after download.
3. If not previously done, set up the connection to CQI database.

- a. On the main menu, click **Options > Database Settings**. Click the CQI tab in the dialog box.

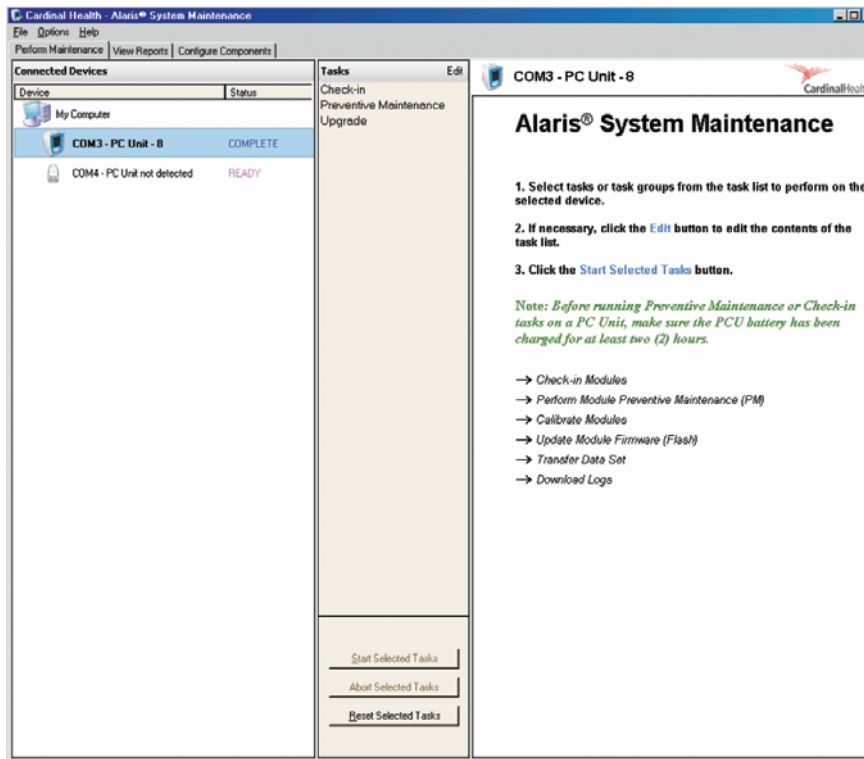


- b. If you need to add a different CQI database, click **Add**.

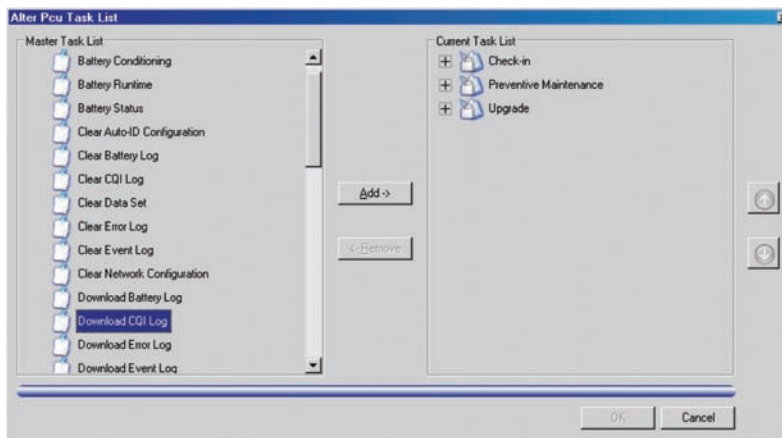


- c. Select the desired SQL Server from the drop-down menu.
- d. From the **Database name** drop-down menu, select the database name for the CQI database.
- e. To verify the connection to the database, click **Test Connection**.
- f. Click **OK**.

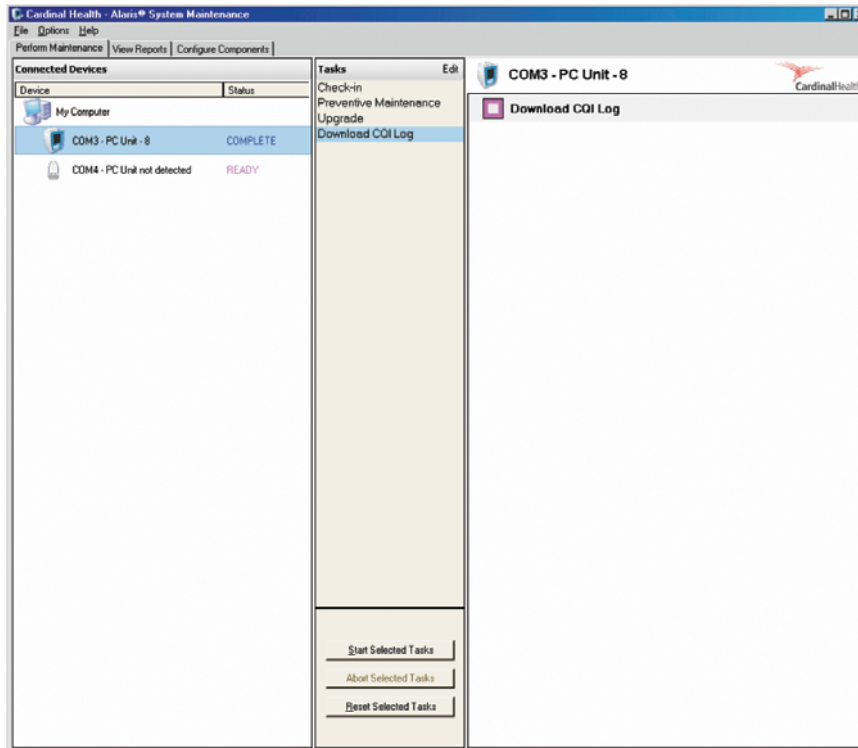
- To perform the download operation, click the desired PC Unit in the Connected Devices pane.



- If the **Download CQI Log** task is not available in the Tasks list, you might want to add it.
 - Click **Edit**.



- b. In the Alter PCU Task List dialog box, click **Download CQI Log**, click **Add**, and then click **OK**.



6. You can now select the **Download CQI Log** task for this PC Unit and then click **Start Selected Task**.

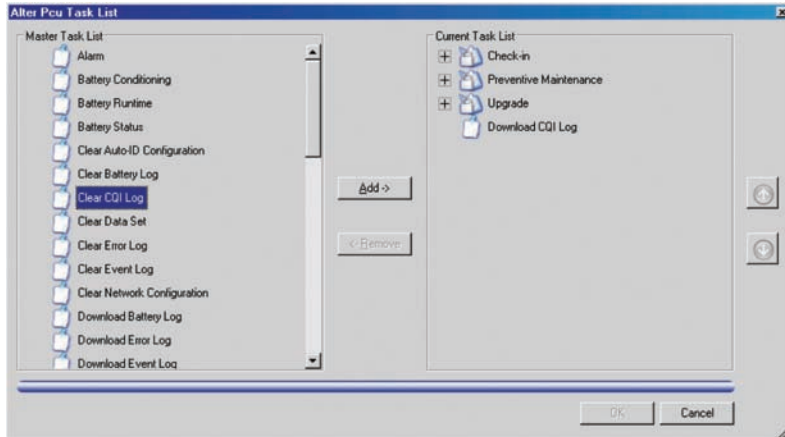
Step 1:

If your facility has multiple sites, facility IDs distinguish one facility from another. This is not a required field but providing an ID for your facility site allows you to either see the CQI Report for a single site or to compare reports between sites. Type in the **Facility ID** or leave the field blank, and then click **Next**.

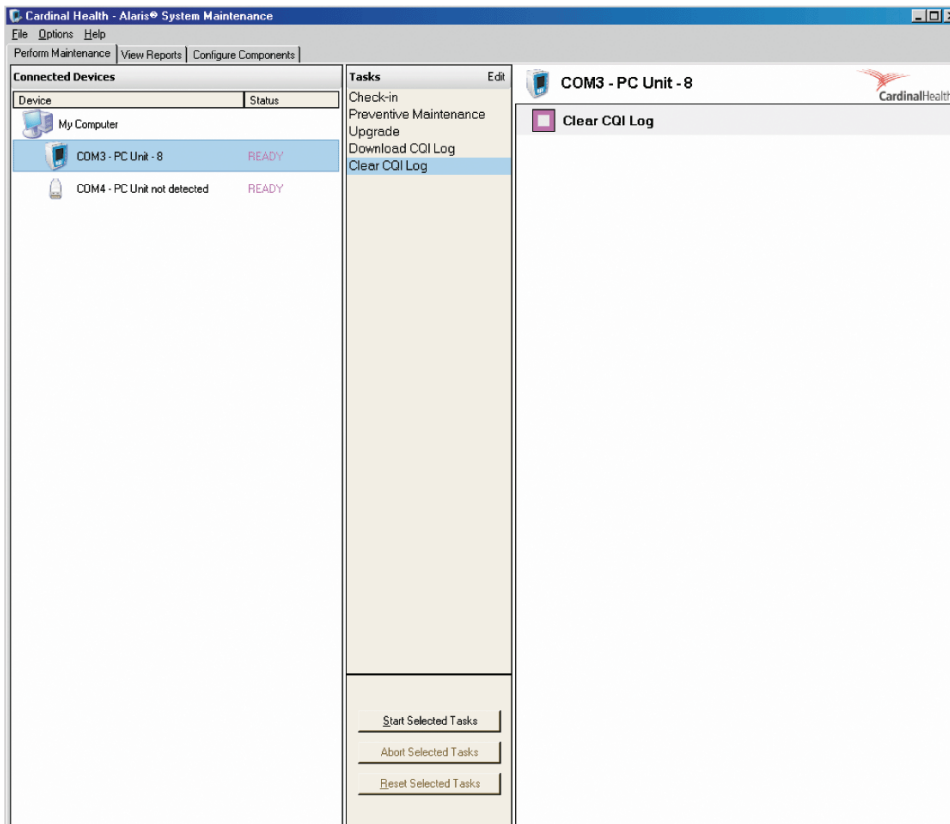
Step 2:

The download operation starts. It usually only takes a few seconds to complete.

7. If you did not select Clear Downloaded CQI Logs, then you should now manually clear the CQI log on the PC unit.
 - a. At the top of the Tasks list, click **Edit**.
 - b. In the Alter PCU Task List dialog box, click **Clear CQI Log**.

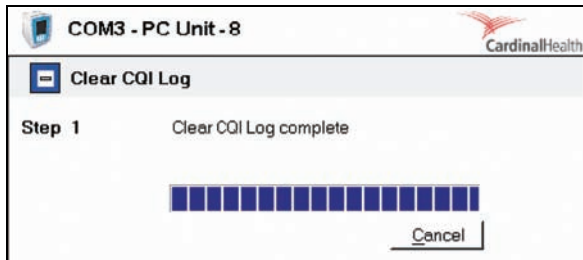


- c. Click **Add** and then click **OK** to save the changes.
The Clear CQI Log task is now available.



8. Click **Clear CQI Log** in the Tasks list and then click **Start Selected Tasks**.

- Click the + icon to the left of **Clear CQI Log** to view the "Clear CQI complete" message.



You have successfully downloaded the CQI log to the CQI database and cleared the log on the PC Unit.

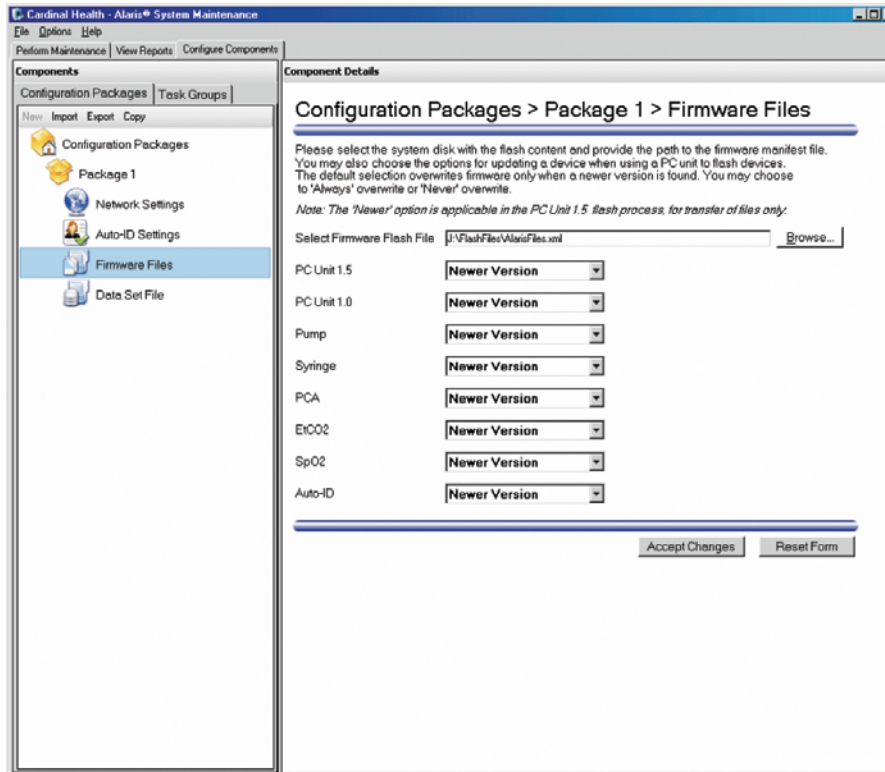
Chapter 8

Flash Firmware

Flash Firmware

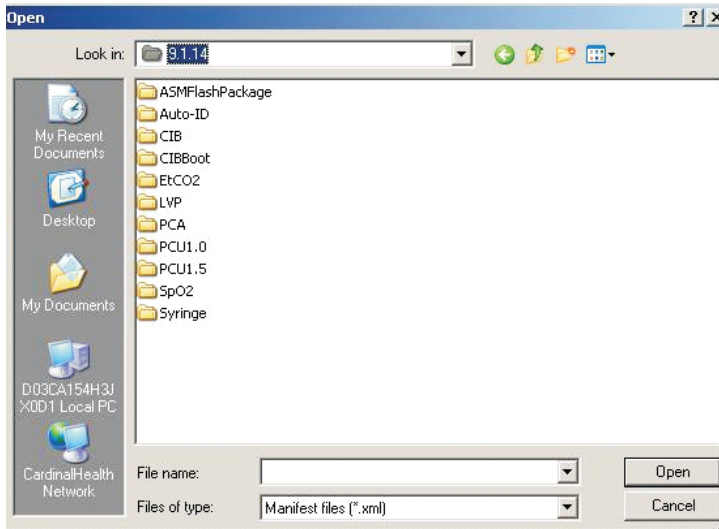
PC Unit 8000 (PCU1.0)

1. Configure the application so that it can access the flash file(s) that will be used to upgrade modules.
 - a. Click the **Configure Components** tab.
 - b. Click the **Configuration Packages** subtab.
 - c. Click **Firmware Files** in the menu.

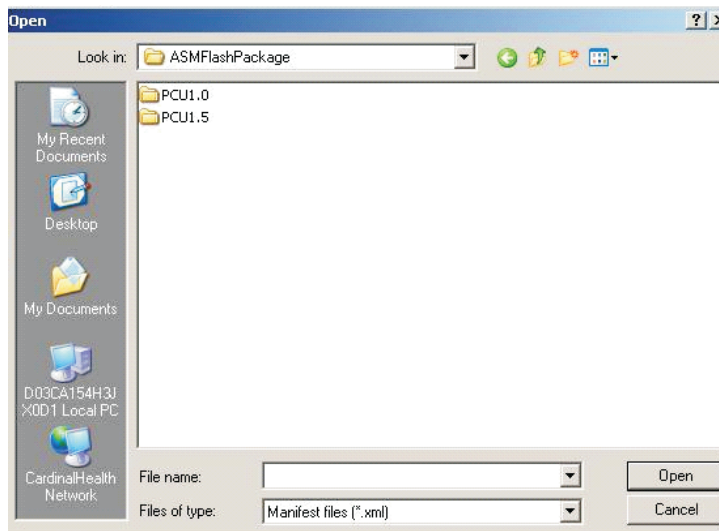


2. If necessary, choose a file for the **Select Firmware Flash File** box by clicking **Browse**.
 - a. Create a folder on your computer and copy the contents of the "Guardrails® Point-of-Care Software, Operating System Software, Guardrails® Software Suite MX" CD into that folder.

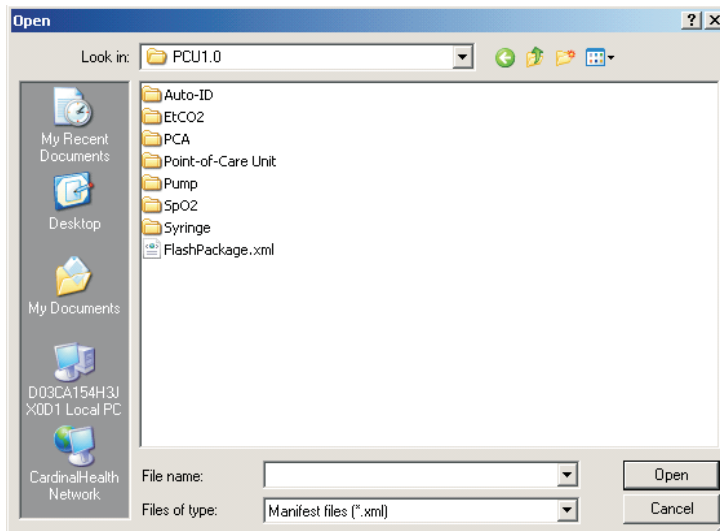
- b. In the **Look in** drop-down menu find and select the folder you created. Double-click **ASMFlashPackage**.



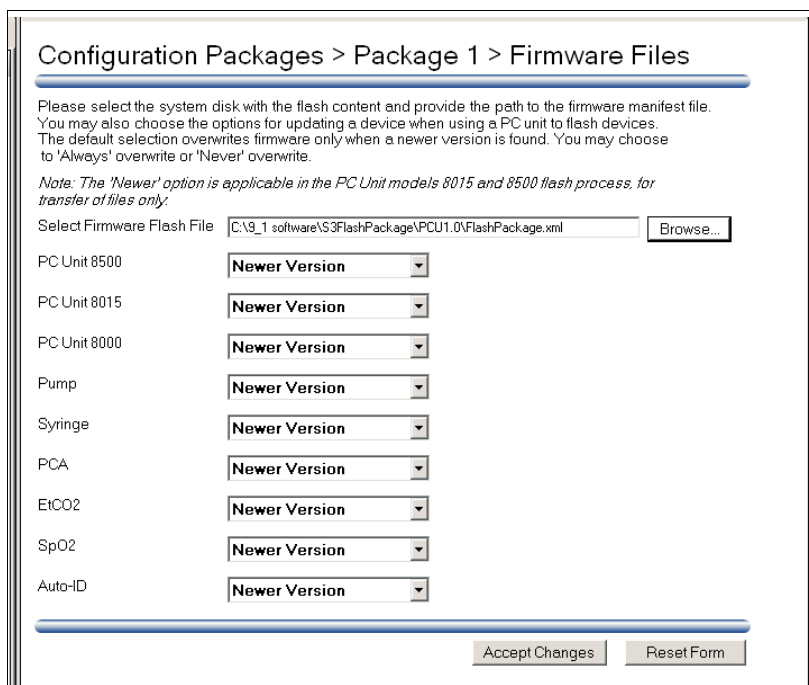
- c. Double-click **PCU1.0**.



- d. Double-click **FlashPackage.xml**.



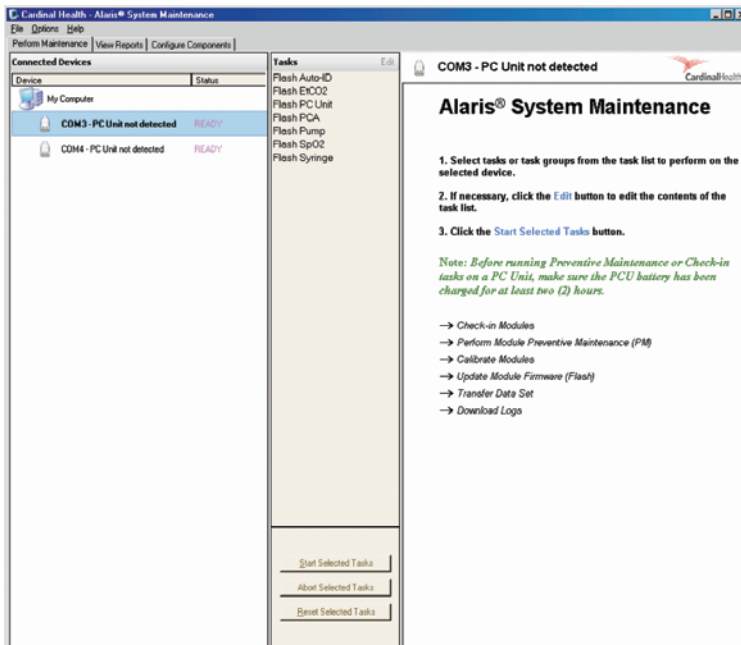
- e. Click **Accept Changes**.



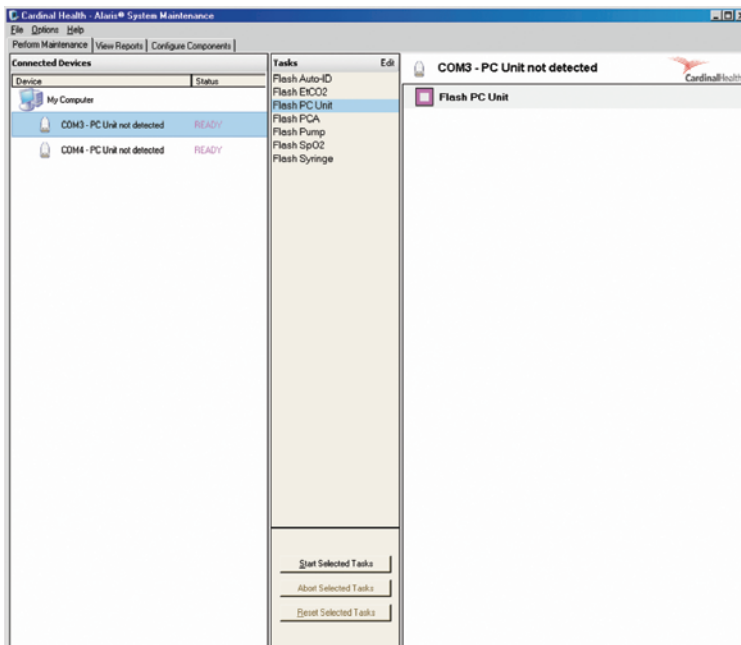
3. From the drop-down menus, select the conditions under which you wish to upgrade each module.
- **Newer Version**—Whenever the user performs an upgrade operation, the module should only be upgraded if the specified firmware version is more recent than the firmware that currently resides on the module.
 - **Always**—The module should always be upgraded with specified firmware file whenever an upgrade operation is performed.
 - **Never**—The module should never be upgraded with specified firmware file whenever an upgrade operation is performed.

4. Click **Perform Maintenance** tab.

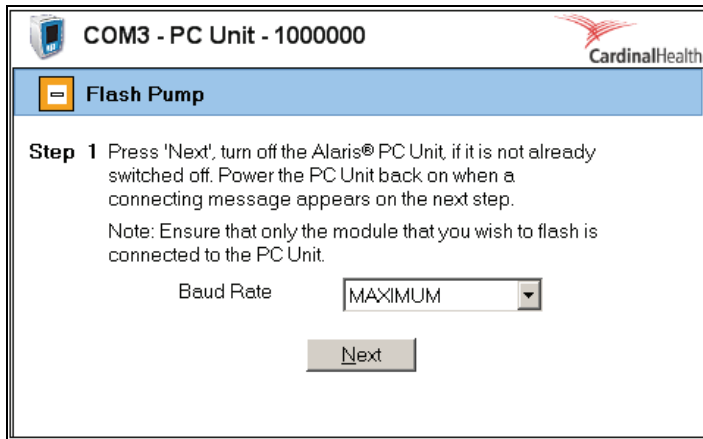
The following window is displayed. It shows available tasks for a PC Unit that is not currently flashed. Note that the task list only includes flash operations.



5. Click **Flash PC unit** and then click **Start Selected Tasks**.

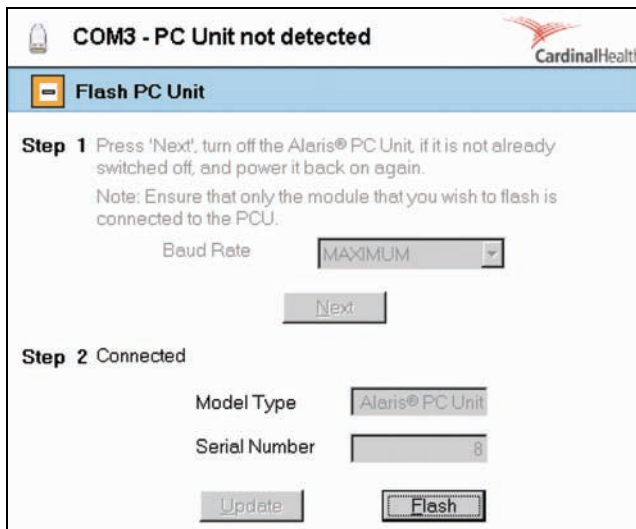


- Follow the instructions on the screen. To turn the PC Unit off, press the **EXIT** soft key twice. To turn the PC Unit back on, press the **SYSTEM ON** key.



- Click **Next**.

If a connection is successfully established, the following window is displayed.

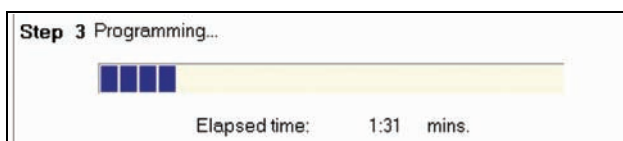


- Click **Flash** to begin flash operation. If this operation fails at the maximum baud rate, try again using a lower baud rate until the process is successful.

The old flash image is erased.



The new flash image is programmed into the device. This step can take several minutes to complete.

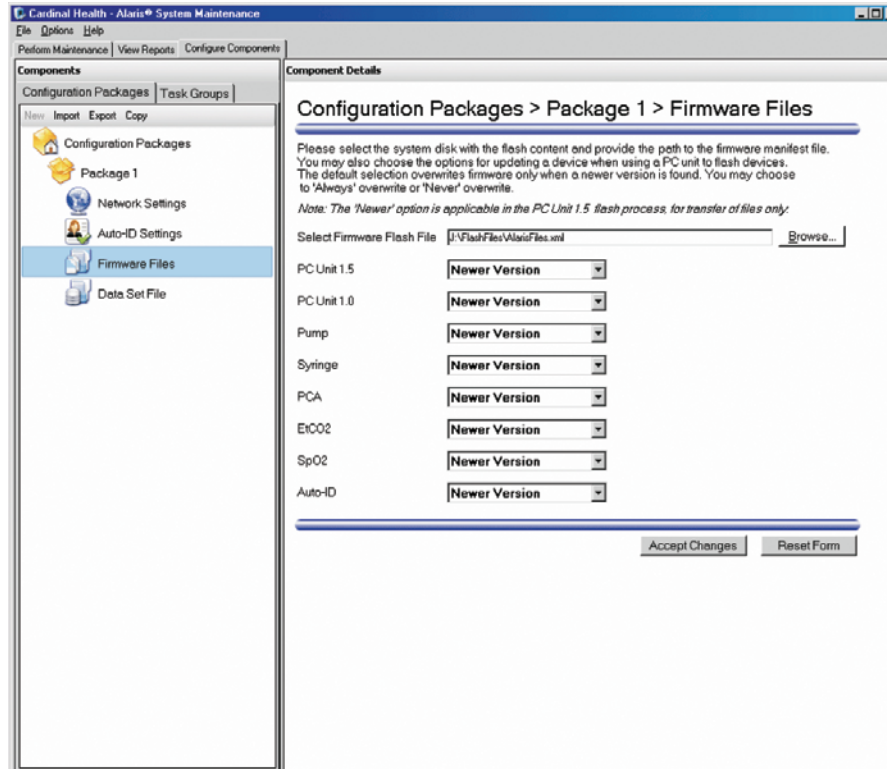


9. When the operation is complete, click **Finish**.

Step 4 Flash completed successfully! Please power down the Alaris® PC unit using the tamper resist switch at the back of the PC Unit and power it back on using the 'System On' key on the unit.
 Press 'Finish' when the PC Unit has been restarted.
 Note: Do not press 'Finish' while another PC Unit is rebooting until the device status is Complete.

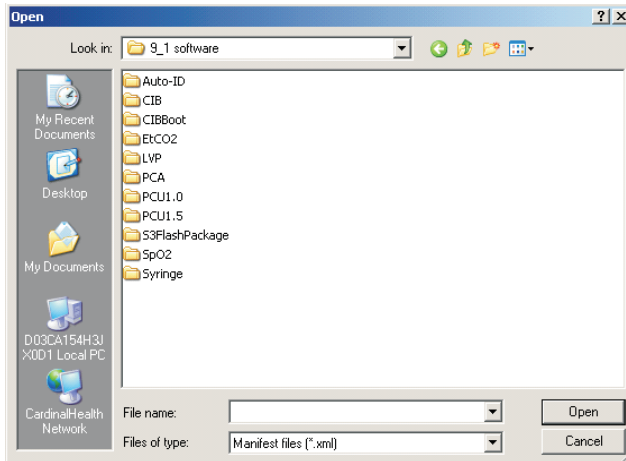
PC Unit 8015 (PCU1.5)

1. Configure the application so that it can access the flash file(s) that will be used to upgrade modules.
 - a. Click the **Configure Components** tab.
 - b. Click the **Configuration Packages** subtab.
 - c. Click **Firmware Files** in the menu.

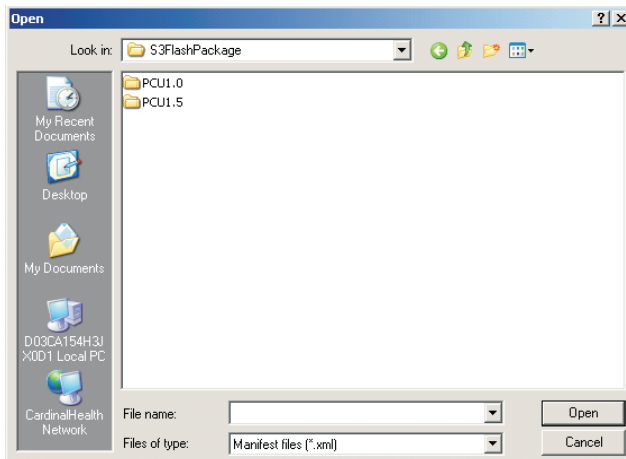


2. If necessary, choose a file for the **Select Firmware Flash File** box by clicking **Browse**.
 - a. Create a folder on your computer and copy the contents of the "Guardrails® Point-of-Care Software, Operating System Software, Guardrails® Software Suite MX" CD into that folder.

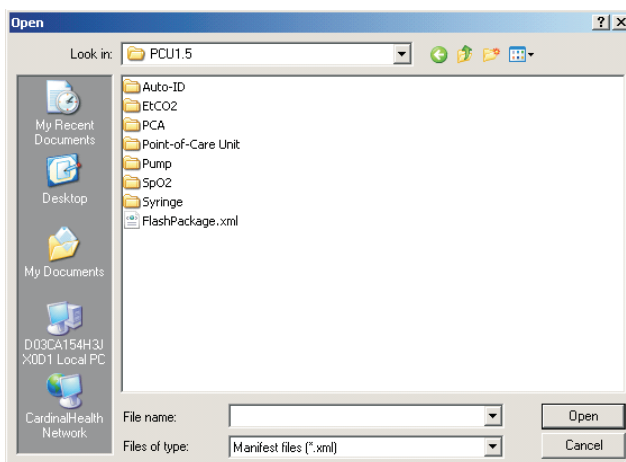
- b. In the **Look in** drop-down menu find and select the folder you created. Double-click **S3FlashPackage**.



- c. Double-click **PCU1.5**.



- d. Double-click **FlashPackage.xml**.



- e. Click **Accept Changes**.

Configuration Packages > Package 1 > Firmware Files

Please select the system disk with the flash content and provide the path to the firmware manifest file. You may also choose the options for updating a device when using a PC unit to flash devices. The default selection overwrites firmware only when a newer version is found. You may choose to 'Always' overwrite or 'Never' overwrite.

Note: The 'Newer' option is applicable in the PC Unit models 8015 and 8500 flash process, for transfer of files only.

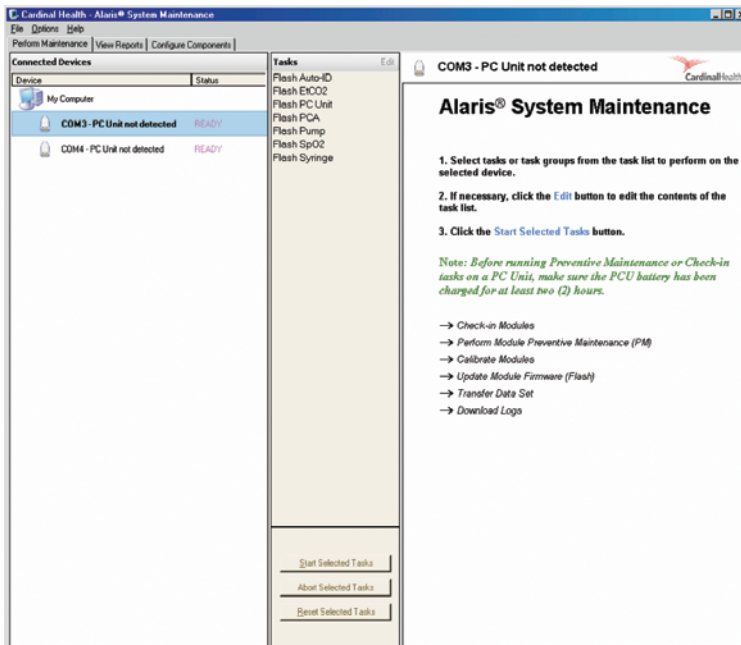
Select Firmware Flash File

PC Unit 8500	<input type="text" value="Newer Version"/>
PC Unit 8015	<input type="text" value="Newer Version"/>
PC Unit 8000	<input type="text" value="Newer Version"/>
Pump	<input type="text" value="Newer Version"/>
Syringe	<input type="text" value="Newer Version"/>
PCA	<input type="text" value="Newer Version"/>
EtCO2	<input type="text" value="Newer Version"/>
SpO2	<input type="text" value="Newer Version"/>
Auto-ID	<input type="text" value="Newer Version"/>

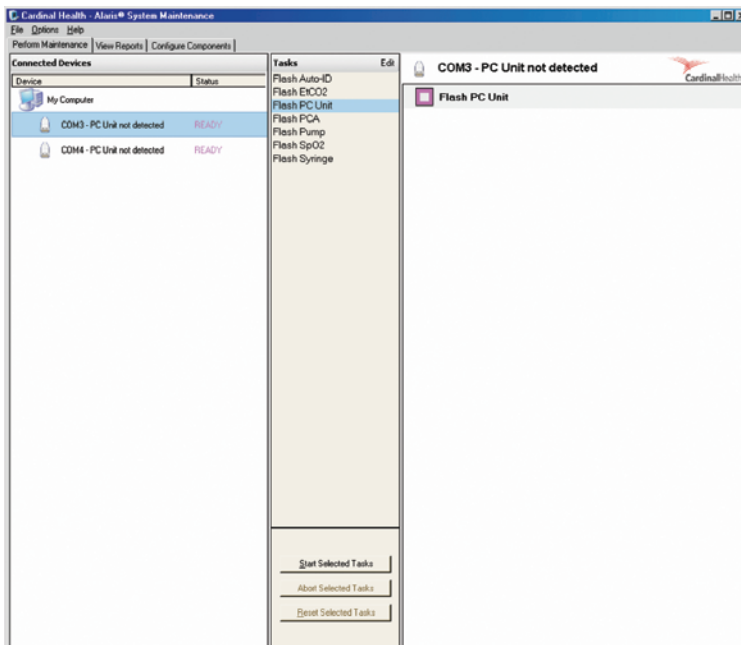
3. From the drop-down menus, select the conditions under which you wish to upgrade each module.
- **Newer Version**—Whenever the user performs an upgrade operation, the module should only be upgraded if the specified firmware version is more recent than the firmware that currently resides on the module.
 - **Always**—The module should always be upgraded with specified firmware file whenever an upgrade operation is performed.
 - **Never**—The module should never be upgraded with specified firmware file whenever an upgrade operation is performed.

4. Click **Perform Maintenance** tab.

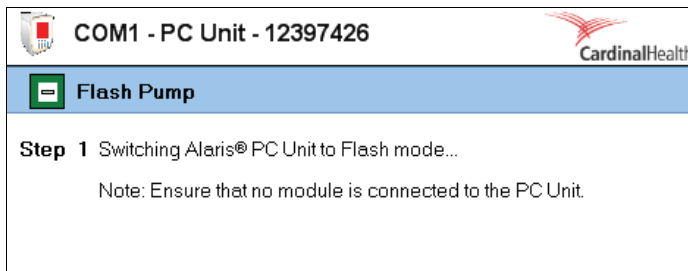
The following window is displayed. It shows available tasks for a PC Unit that is not currently flashed. Note that the task list only includes flash operations.



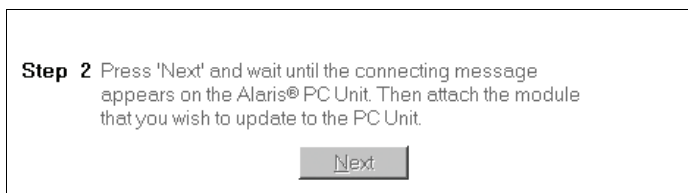
5. Click **Flash PC unit** and then click **Start Selected Tasks**.



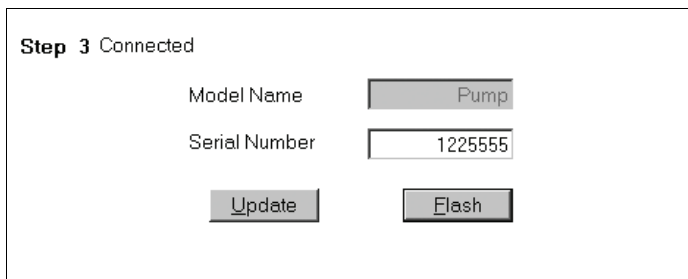
- The software is establishing a connection with the PC Unit and transferring to flash mode. If a connection is successfully established, the following window is displayed.



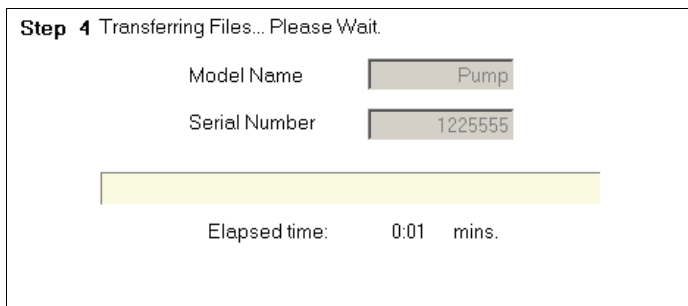
- Click **Next**.



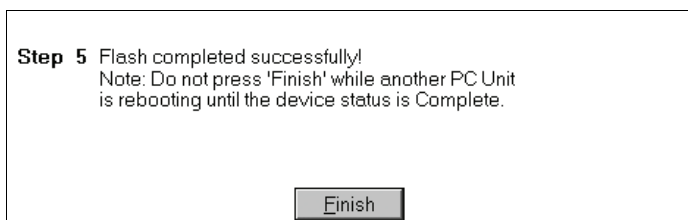
- After the connection is complete, the following connection is displayed. Click **Flash** to begin flashing the device.



- The following window is displayed as the flashing process is executed.



- When the operation is complete, click **Finish**.



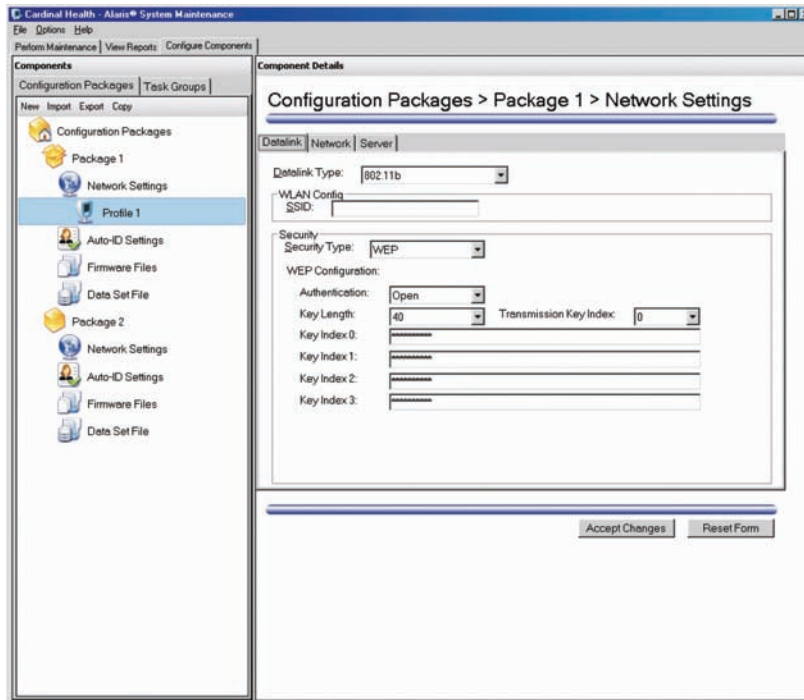
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Chapter 9

Configure Components

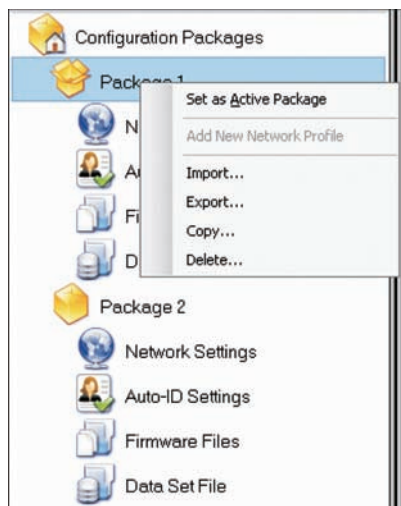
Configuration Packages

Configuration Packages contain the configuration settings that can be transferred to the PC Unit and modules, as well as the configuration information that is used solely on the System Maintenance Software that is running on the personal computer. Multiple Configuration Packages can be defined, as shown below.



One Configuration Package is designated as the active package. The active package is used whenever you transfer configuration information to a PC Unit, or otherwise use the configuration information within the System Maintenance Software. In the example above, Package 1 is the active package, as indicated by the icon of an open box.

To start, click the **Configure Components** tab. To make a package the active package, right-click the package name and then click **Set as Active Package**.



Components of a Configuration Package

Configuration Packages are comprised of the following components:

Network Settings	<p>Used by the Transfer Network Configuration task, the Transfer Network Configuration (Silent) task, and the Network Connectivity task.</p> <p>Network Settings specify the network configuration that is used to connect the PC Unit to Information Server, including:</p> <ul style="list-style-type: none"> • Network security profile • IP addresses • DNS information • Application-level communication protocol information <p>Defining network profiles should only be performed by Network Administrator or other qualified personnel.</p> <p>System Maintenance Software can define and save up to eight Network Profiles on a PC Unit.</p> <p>These profiles are used when a PC Unit connects to a network, allowing a facility to create a large network with multiple access points. If a PC Unit is moved, it cycles through available profiles to find an optimum profile for that location.</p>
Auto-ID Settings	<p>Used by the Transfer Auto-ID Configuration task and the Auto-ID Configuration (Silent) task. Contains configuration information for the Auto-ID Module.</p>
Firmware Files	<p>Used by individual flash tasks that updates the firmware on the PC Unit and modules. Identifies firmware file to use for updating the PC Unit and modules, and contains settings on how to apply updates to specific modules.</p>
Data Set File	<p>Used by the Transfer Data Set task. Specifies the Data Set to transfer to non-networked PC Units.</p>

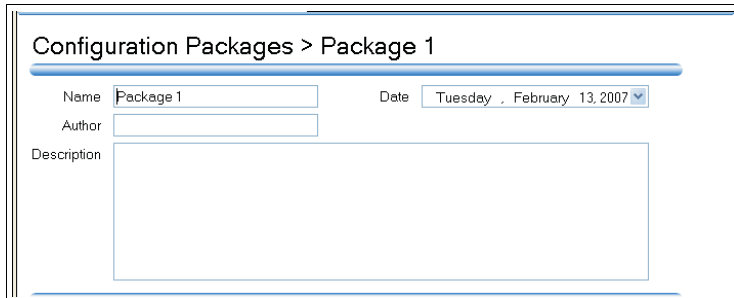
Add a New Configuration Package

To add a new configuration package to the list of available configuration packages:

1. On the main window, click the **Configure Components** tab.
2. In the Components window, click the **Configuration Packages** tab.
3. Click the **Configuration Packages** item and click **New** (or right-click the **Configuration Packages** item and click **Add New Package**).

A new package is added to the list of available packages.

4. Complete the Component Details dialog box:



- a. **Name:** Give the configuration package a unique name.
 - b. **Date:** Select a creation date (defaults to today's date).
 - c. **Author:** Name of person creating configuration package.
 - d. **Description:** A description for configuration package.
5. Click **Accept Changes** to save details or **Reset Form** to discard changes.
 6. Select each configuration package component (Network Settings, Auto-ID Settings, Firmware Files, and Data Set File) and fill in the information.

Delete a Configuration Package

All information in the configuration package will be deleted. If you wish to save the information for later retrieval, export the configuration package before deleting it.

To delete a configuration package:

1. On the main window, click the **Configure Components** tab.
2. In the Components window, click the **Configuration Packages** tab.
3. Right-click the name of the configuration package that you want delete and click **Delete**.

A confirmation dialog box is displayed.

4. To delete the package, click **Yes**.

Export a Configuration Package

You can export a configuration package or components of a configuration package.

1. In the list of Configuration Packages, click the name of a configuration package to highlight it.

2. Click **Export** (or right-click the name of the configuration package and click **Export**).
The Save As dialog box is displayed.
3. Name the export file.
The default name contains the configuration package name and the date and time it was saved.
Exported configuration packages must have a file extension type of `.zip`.

Import a Configuration Package

You can import a configuration package or components of a configuration package. The imported data will overwrite any data already in the selected configuration package. If you do not want to overwrite the data, first create a new configuration package and then import a configuration package into it.

To import a configuration package:

1. In the list of **Configuration Packages**, click the name of a configuration package to highlight it.
2. Click **Import** (or right-click the name of the configuration package and click **Import**).
The Open dialog box is displayed.
3. Select a `.zip` file (or a `.cfg` file if you are importing from a previous version) to import and click **Open**.
Data is imported into the selected configuration package.

NOTE

When a `.cfg` file is imported, any Private Key file, User Certificate file, or CA Certificate file used to create the Network profile being imported must be copied to the same location.

4. Accept any changes in the selected configuration package by clicking **Accept Changes**.

Export or Import Components of a Configuration Package

You can export or import parts of a configuration package by selecting the package at the component level (for example, Auto-ID Module settings) instead of selecting the configuration package name. Then export the configuration package (see *Export a Configuration Package* on page 120) or import it (see).

When importing, you can only import configuration data to overwrite data of the same type in the selected configuration package.

NOTE

The default name of an exported file will include the component type unless it contains a complete configuration package. For example, a complete configuration package might be named

`Home_Package1_3_30_2006_12_00_00_PM.zip`,

while a package named

`Home_Package1_NetworkSettings_3_30_2006_12_00_00_PM.zip`

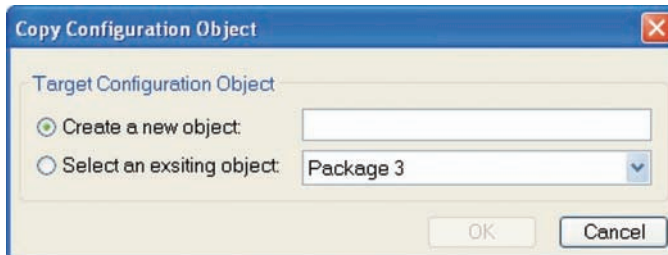
contains only network settings.

Copy a Configuration Package

You can copy the contents of a configuration package into another package. Data in the target configuration object will be overwritten.

1. In the **Configuration Packages** list, click a configuration package name to highlight it.
2. Click **Copy** (or right-click the name of the configuration package and click **Copy**).

The Copy Configuration Object dialog box is displayed.



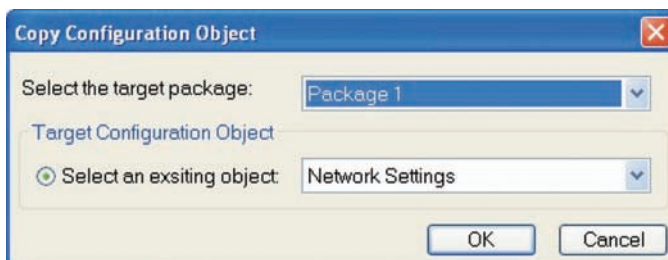
3. Click **Create a new object** and enter a new object name in which to copy the configuration package, or click the **Select an existing object** radio button and then click the object name from the drop-down menu.
4. Click **OK**.
5. Accept changes in target configuration package by clicking **Accept Changes**.

Copy a Component of a Configuration Package

You can copy a component of a configuration package into another package. Data in the target configuration object will be overwritten.

1. In the **Configuration Packages** list, click a component of a configuration package to highlight it.
2. Click **Copy** (or right-click the name of the component and click **Copy**).

The Copy Configuration Object dialog box is displayed.



3. Select the target configuration package from the drop-down menu.
4. Click **OK**.
The configuration component is copied into the target configuration package.
5. Click **Accept Changes** in the target configuration package.

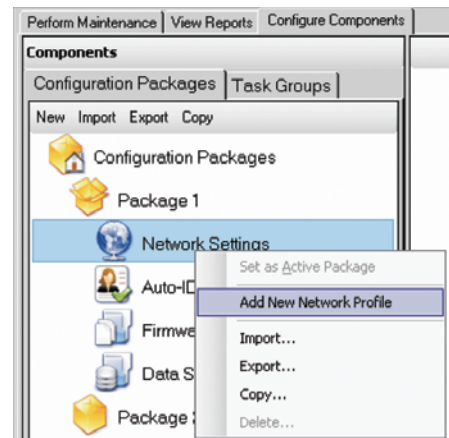
Specify Network Settings

Create a Network Profile

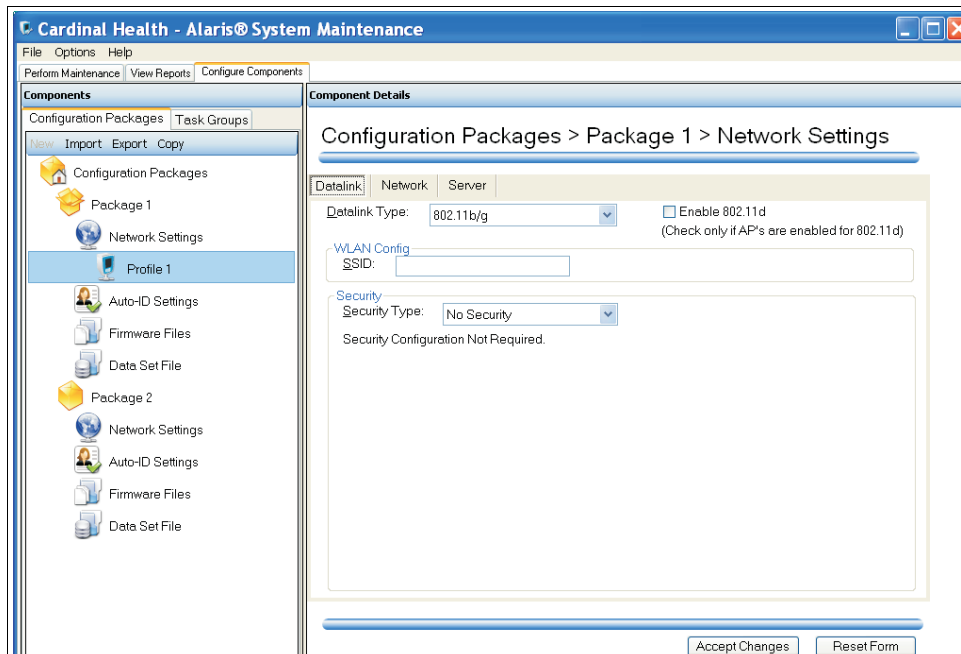
The System Maintenance Software can define and save up to eight network profiles on a PC Unit. These profiles are used when the PC Unit connects to the Information Server. If a PC Unit is moved, it cycles through the available profiles to find the optimum profile to use to connect to the server from that location.

To create a new profile:

1. On the main window, click **Configure Components** tab.
2. In the Components window, click **Configuration Packages** tab.
3. Click **NEW** or right-click **Network Settings** under the desired package.
4. Click **Add New Network Profile**.



The profile is displayed in the Component Details pane.



Datalink Settings

The Datalink tab allows you to specify the wireless security settings for this profile.

Due to incompatibility of the driver for the RF card, WPA2 security will not work with an Alaris® System that is using the LA-4137 Wireless Networker CompactFlash Radio Card from Motorola (sometimes called the Symbol "b" RF card, previously manufactured by Symbol Technologies).

1. Choose the appropriate Datalink Type.

Configuration Packages > Package 1 > Network Settings

Datalink Network Server

Datalink Type: 802.11b/g Enable 802.11d
(Check only if AP's are enabled for 802.11d)

WLAN Config
SSID: 1234

Security
Security Type: No Security
Security Configuration Not Required.

2. Leave 802.11d unchecked (for the U.S. wireless standard); if you want to enable wireless standards for other countries or international country-to-country roaming, check this parameter.

NOTE

If you enable the 802.11d parameter, the international regulatory wireless domain will be activated. The WLAN equipment (wireless controller and access points) have to be configured for 802.11d. If they are not, the wireless PC Units that have network profile 802.11d enabled will not be able to connect to the wireless network.

3. Enter the SSID.
4. Select the appropriate security type.

Configuration Packages > Package 1 > Network Settings

Datalink Network Server

Datalink Type: 802.11a/b/g Enable 802.11d
(Check only if AP's are enabled for 802.11d)

WLAN Config
SSID:

Security
Security Type: No Security
Security Configuration: No Security, WEP, WPA - PSK, WPA - EAP - TLS, WPA - EAP - TTLS, WPA - PEAP, WPA2 - PSK, WPA2 - EAP - TLS

5. Enter the security settings that apply to the selected security type, as follows:

Security Settings for WEP on page 125

Security Settings for WPA or WPA2—with PSK on page 125

Security Settings for WPA or WPA2—with EAP-TLS on page 126

Security Settings for WPA or WPA2—with EAP-TTLS on page 127

Security Settings for WPA or WPA2—with PEAP on page 127

Security Settings

Security Settings for WEP

Security

Security Type: WEP

WEP Configuration:

Authentication: Open

Key Length: 40 Transmission Key Index: 0

Key Index 0:

Key Index 1:

Key Index 2:

Key Index 3:

Accept Changes Reset Form

1. **Authentication**—Select the type of wireless network authentication from the drop-down: Open System or WEP Shared Key.
2. **Key Length**—Select 40-bit or 128-bit encryption type.
3. **Transmission Key Index**—Select one of the four WEP keys to use as the transmission key.
4. **Key Index**—If WEP encryption is enabled, enter the WEP key in the **Key Index 0** field. If needed, up to three additional WEP keys can be entered—one in each of the remaining Key Index fields.

NOTE

Do not click **Accept Changes** until all required entries under the Datalink, Network, and Server tabs are completed.

Security Settings for WPA or WPA2—with PSK

Security

Security Type: WPA-PSK

Encryption:

Type: AES

WPA-PSK:

Passphrase: Hex

Accept Changes Reset Form

1. **Encryption**—Select TKIP (for WPA-PSK) or AES (for WPA2-PSK).
2. **Passphrase**—Enter a passphrase key in ASCII or Hex format.

3. **Hex**—If the passphrase key was entered in Hex format, click the **Hex** checkbox.

NOTE

Do not click **Accept Changes** until all required entries under the Datalink, Network, and Server tabs are completed.

Security Settings for WPA or WPA2—with EAP-TLS

The screenshot shows a configuration window titled "Security". The "Security Type" dropdown is set to "WPA2-EAP-TLS". Under "Encryption", the "Type" dropdown is set to "AES". The "WPA2 EAP TLS" section includes an "Identity" text field. Below it are "Private Key" and "User Certificate" sections, each with a "File" selection button, a "Format" dropdown menu, and a "Passphrase" text field. The "CA Certificate" section has a "Validate Server Certificate" checkbox and a "File" selection button. At the bottom of the window are "Accept Changes" and "Reset Form" buttons.

1. **Encryption**—Select TKIP (for WPA-EAP-TLS) or AES (for WPA2-EAP-TLS).
2. **Identity**—Enter the identity name.
3. **Private Key**—Import the private key from a file.
4. **Format** (for Private Key)—Select the format: BER, DER, or PKCS12.
5. **Passphrase** (for Private Key)—If the PKCS12 format is selected, enter the passphrase key.
6. **User Certificate**—Import the user certificate from a file.
7. **Format** (for User Certificate)—Select the format: BER, DER, or PKCS12.
8. **Passphrase** (for User Certificate)—If the PKCS12 format is selected, enter the passphrase key.
9. **Validate Server Certificate** (optional)—If server validation is required on the network, click the **Validate Server Certificate** box to enable server validation.
10. **CA. Certificate**—If **Validate Server Certificate** is enabled, import the CA certificate from a file.
11. **Format** (for CA. Certificate)—Select the format: BER, DER, or PKCS7.

NOTE

Do not click **Accept Changes** until all required entries under the Datalink, Network, and Server tabs are completed.

Security Settings for WPA or WPA2—with EAP-TTLS

The screenshot shows a configuration window titled "Security" with the following fields and options:

- Security Type:** WPA2 - EAP - TTLS (dropdown menu)
- Encryption:**
 - Type:** AES (dropdown menu)
- WPA2 EAP TTLS:**
 - Identity:** (text input field)
 - Validate Server Certificate
- CA. Certificate:**
 - File: (text input field)
 - Format: (dropdown menu)
- Inner Authentication:**
 - Type:** (dropdown menu)
 - User Name:** (text input field)
 - Password:** (text input field)

At the bottom of the window are two buttons: "Accept Changes" and "Reset Form".

1. **Encryption**—Select TKIP (for WPA-EAP-TTLS) or AES (for WPA2-EAP-TTLS).
2. **Identity**—Enter the identity name.
3. **Validate Server Certificate** (optional)—If server validation is required on the network, click the **Validate Server Certificate** box to enable server validation.
4. **CA. Certificate**—If **Validate Server Certificate** is enabled, import the CA certificate from a file.
5. **Format** (for CA. Certificate)—Select the format: BER, DER, or PKCS7.
6. **Inner Authentication Type**—Select MSCHAPv2 or PAP.
7. **User Name**—Enter a valid user name (for example, "jsmith"). Most of the time the user name and identity are the same.
8. **Password**—Enter the correct user name password.

NOTE

Do not click **Accept Changes** until all required entries under the Datalink, Network, and Server tabs are completed.

Security Settings for WPA or WPA2—with PEAP

The screenshot shows a configuration window titled "Security" with the following fields and options:

- Security Type:** WPA2 - PEAP (dropdown menu)
- Encryption:**
 - Type:** AES (dropdown menu)
- WPA2 EAP PEAP:**
 - Identity:** (text input field)
 - Validate Server Certificate
- CA. Certificate:**
 - File: (text input field)
 - Format: (dropdown menu)
- Inner Authentication:**
 - Type:** (dropdown menu)
 - User Name:** (text input field)
 - Password:** (text input field)

At the bottom of the window are two buttons: "Accept Changes" and "Reset Form".

1. **Encryption**—Select TKIP (for WPA-PEAP) or AES (for WPA2-PEAP).
2. **Identity**—Enter the identity name.
3. **Validate Server Certificate** (optional)—If server validation is required on the network, click the **Validate Server Certificate** box to enable server validation.
4. **CA. Certificate**—If **Validate Server Certificate** is enabled, import the CA certificate from a file.
5. **Format** (for CA. Certificate)—Select the format: BER, DER, or PKCS7.
6. **Inner Authentication Type**—Select MSCHAPv2.
7. **User Name**—Enter a valid user name (for example, "jsmith"). Most of the time the user name and identity are the same.
8. **Password**—Enter the correct user name password.

NOTE

Do not click **Accept Changes** until all required entries under the Datalink, Network, and Server tabs are completed.

Network Settings

The **Network** tab allows you to specify the Internet Protocol (IP) information for connecting to the wireless network. In the drop-down menu, choose one of the following options:

- **Obtain an IP Address Automatically.** The PC Unit will automatically receive an IP address from the Dynamic Host Configuration Protocol (DHCP) server on the network. Note that the system also automatically configures **Obtain DNS Server Address Automatically** as the default setting (the Preferred DNS Server and Alternate DNS Server will automatically receive an IP address from the DHCP server on the network).
- **Use Static IP Address.** The PC Unit will be assigned a static IP address. If you choose this option, you must enter an IP Address, Subnet, and Default Gateway. Note that the system automatically configures **Use Static DNS IP Address** as the default setting.
 - **Static IP Address**—The IP address has to be entered or assigned manually on each PC Unit.

- **IP Address range**—The auto-assign option helps ensure that multiple PC Units are not given the same IP address. Enabling auto-assign provides a range of IP addresses to use for a subnet and the software automatically assigns an IP address to a connected PC Unit when it is updated on that subnet. The software maintains a table of IP addresses for this purpose.

When a PC Unit is updated on the network, the System Maintenance Software confirms that the PC Unit has an IP address assigned to it for each auto-assigned subnet. If an IP address is not assigned for a subnet, the System Maintenance Software applies the next available IP address in the configured range and assigns it to the PC Unit.

NOTE

Do not click **Accept Changes** until all required entries under the Datalink, Network, and Server tabs are completed.

Server Settings

The screenshot shows a configuration window with three tabs: "Datalink", "Network", and "Server". The "Server" tab is selected. The form contains the following fields:

- Server Type: Dcmp (dropdown menu)
- Alaris® Server Information:
 - AES Encryption: 128 (dropdown menu)
 - Encryption Key: [text input field] [Import... button]
 - Host Name: [text input field]
 - Server Port: 1 (text input field)

At the bottom of the window are two buttons: "Accept Changes" and "Reset Form".

The Server tab allows you to specify the application-level communication protocol settings that will be used between the PC Unit and the Alaris® Server.

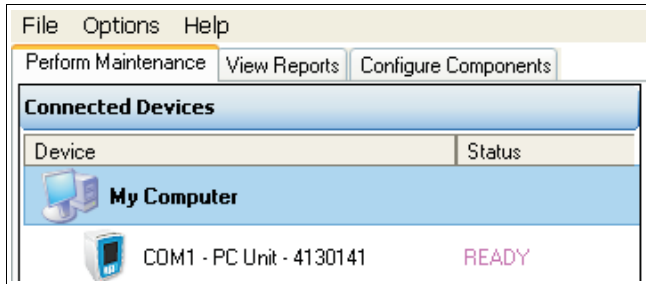
1. **Server Type**—Select **Dcmp** from the drop-down menu.
2. **AES Encryption**—Select an encryption length of 128-bit.
3. **Encryption Key**—If you do not use a server information file exported from the Alaris® Server, enter an encryption key with the correct number of hexadecimal characters (32 characters for 128-bit). When entered, the encryption key is displayed as asterisks (*). If the key is imported from the Alaris® Server file, it is always displayed as asterisks.
4. **Host Name**—Enter the fully-qualified domain name (FQDN) or the IP address of the Alaris® Server.
5. **Server Port**—Enter the server port for the Alaris® Server that the device will be connecting to (Example: 3613).
6. Click **Accept Changes** to save the network profile configuration.

Wireless Network Setting on PC Unit—Disable, Enable, View Status

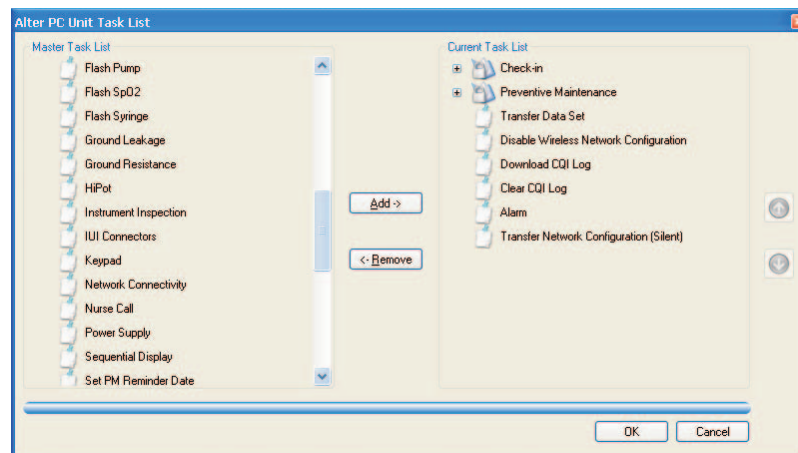
Enable Wireless Network

If your PC Unit will be operating in a wireless environment and the wireless network was disabled using the Maintenance Software, perform the following procedure to enable the connection:

1. Click the **Perform Maintenance** tab.

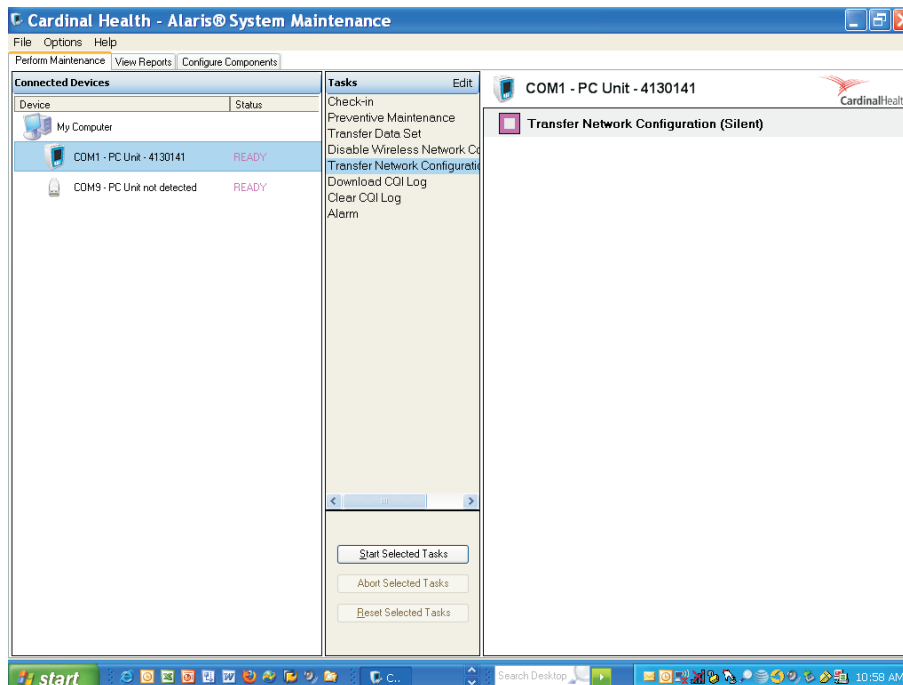


2. In the Connected Devices pane, click the PC Unit that is to have the Wireless Network Configuration enabled.
3. If **Transfer Network Configuration (Silent)** is in the Current Task List, proceed to step 6; otherwise, continue with the next step.
4. Click **Edit** in the Tasks list.
5. Click **Transfer Network Configuration (Silent)** in the Tasks list and click **Add**.

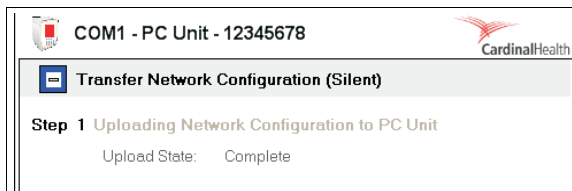


6. Click **OK** to save the change.

- Click **Transfer Network Configuration (Silent)** in the Tasks list.



- Click **Start Selected Tasks** to enable the wireless network configuration on the PC Unit.
- Click the + icon to the left of **Transfer Network Configuration (Silent)** to view when "Uploading Network Configuration to the PC Unit" is complete.



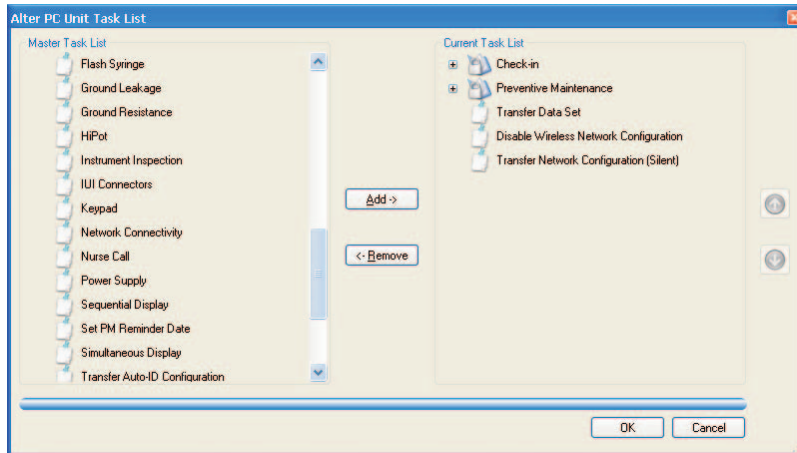
- If desired, verify that the network settings have been transferred. Refer to the System Options, Network Status section in the Alaris® System DFU and use password **32221** to view the wireless status.

Disable Wireless Network

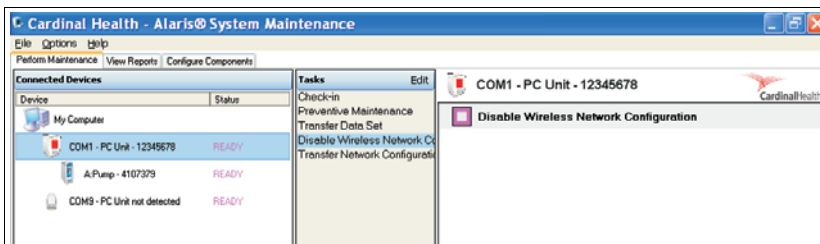
If your PC Unit will be operating in a non-wireless environment, you can disable the wireless network. The connection can be temporarily disabled by changing the System Options setting on the PC Unit or the Maintenance Software can be used to permanently disable the wireless network configuration setting. To permanently disable the wireless network on the PC Unit, perform the following procedure:

- In the Connected Devices pane, click the PC Unit that is to have the Wireless Network Configuration disabled.
- If **Disable Wireless Network Configuration** is in the Current Task List, proceed to step 6; otherwise, continue with the next step.
- Click **Edit** in the Tasks list.

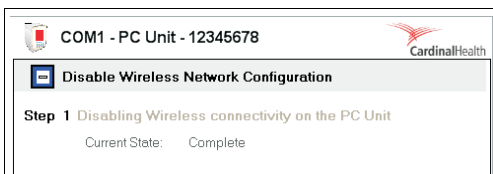
4. Click **Disable Wireless Network Configuration** in the Tasks list. Click **Add** to add it to the Current Task List.



5. Click **OK** to save the change.
6. Click **Disable Wireless Network Configuration** in the Tasks list.



7. Click **Start Selected Tasks** to disable the wireless network configuration on the PC Unit.
8. Click the icon to the left of **Disable Wireless Network Configuration** to view when "Disabling Network connectivity on the PC Unit" is complete.



9. If desired, verify that the wireless network has been disabled.
 - a. Refer to the System Options, Wireless Connection section in the Alaris® System DFU.
 - b. If the **Wireless Connection** soft key is grayed out, wireless communication has been disabled.

Specify Auto-ID Module Settings

1. Click **Configure Components** tab.
2. Click **Configuration Packages** tab.
3. Click **Auto-ID Settings** under the desired package.
4. Use the Component Details pane on the right to define the Auto-ID Module configuration.

Formatting Rules

The following sections identify and explain the Auto-ID settings.

ID Type:

The **ID Type** displays the current rule for identifying Patient and Clinician fields on the bar code labels. Click the **Patient** or **Clinician** button to edit or define its related **ID Format** and **ID Locators**.

NOTE

The screens illustrated in the following sections have **Patient** selected as the **ID Type**. The options for **Clinician** are the same as for **Patient** but the settings must be different.

ID Format:

The **ID Format** is used to select the rule for parsing an ID and to enter corresponding data. Data fields change depending on the rule selected. The **Description** and **Example** fields display information that explains the selected rule.

- **None**—No rule is associated with the **Selected Type**. Entries in the **Data** and **ID Locators** fields are not needed.

Component Details

Configuration Packages > Package 1 > Auto-ID Settings

ID Type

Selected Type	Current Rule
<input checked="" type="radio"/> Patient	Entire String
<input type="radio"/> Clinician	None

ID Format

Rule: None Entire String Field Delimited Character Length
 Tag Length Tagged HL7

Data:

Description:

Example:

ID Locators

Label Min Length: ID Min Length:
Label Max Length: ID Max Length:
Label Prefix:
Label Suffix:

Accept Changes Reset Form

- **Entire String**—The entire string is the ID with no delimiters or field tags. Entries in the **Data** fields are not required.

Component Details

Configuration Packages > Package 1 > Auto-ID Settings

ID Type

Selected Type	Current Rule
<input checked="" type="radio"/> Patient	Entire String
<input type="radio"/> Clinician	None

ID Format

Rule: None Entire String Field Delimited Character Length

Tag Length Tagged HL7

Data:

Description: The entire string is the ID with no delimiters or field tags.

Example: 456123

- **Field Delimited**—The ID is embedded in a delimited string and is Xth field. A **Delimiter** entry in the **Data** field is optional. The **Field** entry must be a numeric value.

Component Details

Configuration Packages > Package 1 > Auto-ID Settings

ID Type

Selected Type	Current Rule
<input checked="" type="radio"/> Patient	Entire String
<input type="radio"/> Clinician	None

ID Format

Rule: None Entire String Field Delimited Character Length

Tag Length Tagged HL7

Data: Delimiter Field

Description: The ID is embedded in a delimited string and is the Xth field

Example: DOE,JIM,456123,04/04/40 – 2nd field (zero based), comma delimited

- Character Length**—The ID starts at the Xth character and is Y characters in length. **Index** (Xth character) and **Length** (Y characters) must be entered in the **Data** fields. The **Length** value must be within the minimum and maximum ID length range. The **Index** value plus the **Length** value must not be greater than the **Label Max Length**. (*ID Locators*: on page 136 for explanations of the minimum and maximum length values.)

Component Details

Configuration Packages > Package 1 > Auto-ID Settings

ID Type

Selected Type	Current Rule
<input checked="" type="radio"/> Patient	Entire String
<input type="radio"/> Clinician	None

ID Format

Rule: None Entire String Field Delimited Character Length

Tag Length Tagged HL7

Data: _____ Index:

Length:

Description: The ID will start at the Xth character and is Y characters in length

Example: 04/04/40456123JIMDOE – 8th character (zero based), 6 characters long

- Tag Length**—The ID is prefixed with a tag and is X characters long. **Tag** characters and **Length** (X characters) must be entered in the **Data** fields. The **Tag** field has a 32-character limit. The **Length** entry must be a numeric value in the range of 1 to 32767.

Component Details

Configuration Packages > Package 1 > Auto-ID Settings

ID Type

Selected Type	Current Rule
<input checked="" type="radio"/> Patient	Entire String
<input type="radio"/> Clinician	None

ID Format

Rule: None Entire String Field Delimited Character Length

Tag Length Tagged HL7

Data: Tag Length:

Description: The ID is prefixed with a tag and is X characters long

Example: LNDOEFNJIMID456123DOB04/04/40 – Beginning tag 'ID', 6 characters long

- **Tagged**—The ID has a **Beginning Tag** and an **Ending Tag**, which must be entered in the **Data** fields. The **Data** fields have a 32-character limit.

Component Details

Configuration Packages > Package 1 > Auto-ID Settings

ID Type

Selected Type	Current Rule
<input checked="" type="radio"/> Patient	Entire String
<input type="radio"/> Clinician	None

ID Format

Rule: None Entire String Field Delimited Character Length
 Tag Length Tagged HL7

Data: Beginning Tag
Ending Tag

Description: The ID has a beginning tag and an ending tag
Example: <DOB>04/04/40</DOB><ID>456123</ID> – Beginning tag '<ID>', Ending Tag

- **HL7**—The ID is represented in an HL7 format. **Tag** and **Field** values must be entered in the **Data** fields. The **Tag** field has a 32-character limit. The **Field** entry must be a numeric value in the range of 1 to 32767. A **Delimiter** entry in the **Data** field is optional.

Component Details

Configuration Packages > Package 1 > Auto-ID Settings

ID Type

Selected Type	Current Rule
<input checked="" type="radio"/> Patient	Entire String
<input type="radio"/> Clinician	None

ID Format

Rule: None Entire String Field Delimited Character Length
 Tag Length Tagged HL7

Data: Tag Field
Delimiter

Description: The ID is represented in an HL7 format
Example: ID|1|22|456123 – Beginning tag 'ID', delimiter '|', 3rd field (zero based)

ID Locators:

Before attempting to extract data from a bar code label, the label data is subjected to validation. The **ID Locators** fields control that validation.

- **Label Min Length**—The minimum number of characters displayed on a label must be at least 1.
- **Label Max Length**—The maximum number of characters displayed on a label must be equal to or greater than the minimum label length.
- **ID Min Length**—The minimum number of characters that make up an ID must be at least 1.
- **ID Max Length**—The maximum number of characters that make up an ID must be equal to or greater than the minimum ID length.
- **Label Prefix**—An ID prefix has a 200-character limit and is optional.

- **Label Suffix**—An ID suffix has a 200-character limit and is optional.

The screenshot shows a configuration window titled "ID Locators". It contains four spinners: "Label Min Length" (set to 1), "Label Max Length" (set to 1), "ID Min Length" (set to 1), and "ID Max Length" (set to 1). Below the spinners are two text input fields: "Label Prefix" and "Label Suffix". At the bottom right of the window are two buttons: "Accept Changes" and "Reset Form".

Specify Firmware Files

1. Use the Files dialog box to identify the Release Manifest File for upload to the PC Unit and modules.
 - a. Firmware Flash File: Use **Browse** to find the correct XML file to be used to update the Alaris® System.
 - b. Device drop-down menu: Choose options from the drop-down menu next to each applicable device, to select what should happen when a PC Unit or module is flashed with contents of the Release Manifest File.
 - **Always:** Always overwrite the contents of the device with content from the Release Manifest File.
 - **Newer Version:** Only overwrites the contents of the device when the Release Manifest File contains a newer version than the one that already existing on the device.
 - **Never:** Do not update the device with the Release Manifest File.
2. Click **Accept Changes** to save edits.

Specify a Data Set File

Use the Data Set File dialog box to select the **Guardrails® data set.GRE** file type for transfer to the PC Unit.

1. Use **Browse** to select the file on the PC workstation.
2. If you click **Always Overwrite Dataset**, whenever you transfer a Data Set to a PC Unit, the old Data Set is overwritten, even if the Data Sets are exactly the same. If you do not click **Always Overwrite Dataset**, the old Data Set is only overwritten if CRC codes for the two Data Sets are different.
3. Click **Accept Changes**.

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Chapter 10

Calibration

Introduction

Calibration can be performed on connected modules by selecting from a list of automated tasks. The calibration task group applies to the Pump Module, PCA Module, Syringe Module, and EtCO₂ Module.

To perform calibration, see the procedure in this section that applies to the module being tested:

Pump Module Calibration on page 141

PCA Module Calibration on page 148

Syringe Module Calibration on page 154

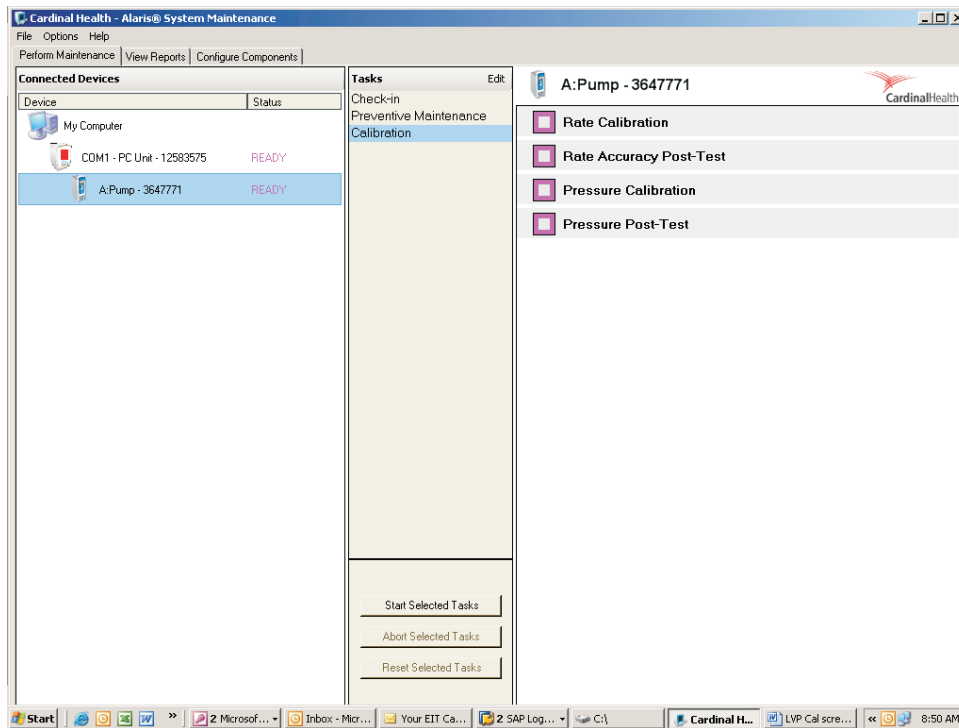
EtCO₂ Module Calibration on page 161

Pump Module Calibration

Test Equipment

Test Equipment	Manufacturer	Model/Part Number	Application
gauge, pressure, digital (peak hold)	Either of the following: <ul style="list-style-type: none"> Heise (www.heise.com) Ashcroft (www.ashcroft.com) 	<ul style="list-style-type: none"> PTE-1 (accuracy from 0.1 to 0.025% span) 2089, 2086, or 2084 (accuracy from $\pm 0.05\%$, 0.10%, or 0.25% of span) or an equivalent gauge with: <ul style="list-style-type: none"> (a) unit of measure in mmHg (b) accuracy of $\pm 1\%$ (c) range of 0-1500 mmHg 	pressure test
hemostat	N/A	N/A	fluid-side occlusion test
IV pole, standard	CareFusion	903-0336 or equivalent	rate, pressure, and occlusion tests
IV sets <ul style="list-style-type: none"> IV set, standard, without check valve IV set, calibration IV set, calibration 	<ul style="list-style-type: none"> CareFusion CareFusion CareFusion 	<ul style="list-style-type: none"> 2210-0500 8100-RCS 8100-PCS 	rate and pressure test
IV solution container (bag preferred)	N/A	N/A	rate and pressure test
luer lock, female, 1/8"-27 NPT	Cole-Parmer (www.coleparmer.com)	K-45503-78 or equivalent	pressure test
reducer, female, 1/4" NPT (F) x 1/8" NPT (F)	Cole-Parmer (www.coleparmer.com)	K-06349-91 or equivalent	pressure test
tubing, silicone	CareFusion	303109 or equivalent	pressure test
T-fitting	CareFusion	303815 or equivalent	pressure test
valve, 3-way	CareFusion	97555 or equivalent	pressure test

1. Click **Perform Maintenance** tab.
2. In the Connected Devices pane, click the Pump Module to be tested.
3. Click **Calibration** in the Tasks list (double-click to immediately begin first task).



4. Click the calibration task that needs to be performed and follow the instructions displayed on the screen.

NOTE

See *Pump Module Tasks* on page 175 for an explanation of each task.

5. Rate Calibration:

Rate Calibration

Step 1 Load a characterized set and prime it with distilled water.
Enter the expected volume provided by the characterization label on the packaging of the set.
Passing values are 11.640 to 12.360 mL.

Expected Volume : grams

- a. Set up the Pump Module for Rate Calibration (see *Rate Accuracy Test Setup* on page 52).
- b. Enter the Expected Volume identified on the label attached to the characterized set.
- c. Click **Next** to prime the set.

Step 2 Priming the line into the container

VTBI	4 ml
Rate	500 ml/h
Elapsed Time	6 secs
Infused Volume	0.5 ml

Step 3 Balance the scale to the zero mark.

Step 4 Rate measure in progress

VTBI	12 ml
Rate	500 ml/h
Elapsed Time	13 secs
Infused Volume	1.5 ml

Ensure that the distilled water is dripping into the cup and that the tubing is not resting on the scale.

Step 5 Enter actual weight reading from scale. Passing values are 11.59 to 12.41 grams.

Actual weight : grams

Step 6 Summary of Test Results

Pump Module Test Result:	Failed
VPMR:	175.3
VTBI:	12 ml
Rate:	500 ml/h
Expected Weight	12.000 grams
Actual Weight	11.51 grams
Acceptable Error	+/-3.400%
Actual Error	-4.0833%

Pump Module Calibrated

Old VPMR	175.3
New VPMR	168.1

6. Rate Accuracy Post-Test:

Set up the Pump Module for Rate Calibration (see *Rate Accuracy Test Setup* on page 52).

Rate Accuracy Post-Test

Step 1 Load a characterized set and prime it with distilled water. Enter the expected volume provided by the characterization label on the packaging of the set. Passing values are 11.640 to 12.360 mL.

Expected Volume : grams

The **Expected Volume** is identified on the label attached to the characterized set.


Step 2 Priming the line into the container

VTBI	4 ml
Rate	500 ml/h
Elapsed Time	6 secs
Infused Volume	0.5 ml

Step 3 Balance the scale to the zero mark.

Step 4 Rate measure in progress

VTBI	12 ml
Rate	500 ml/h
Elapsed Time	13 secs
Infused Volume	1.5 ml



Ensure that the distilled water is dripping into the cup and that the tubing is not resting on the scale.

Step 5 Enter actual weight reading from scale. Passing values are 11.59 to 12.41 grams.

Actual weight: grams

Step 6 Summary of Test Results

Pump Module Test Result	Passed
VPMP:	168.1
VTBI:	12 ml
Rate:	500 ml/h
Expected Weight	12.000 grams
Actual Weight	11.95 grams
Acceptable Error	+/-3.400%
Actual Error	-0.4167%

If the Pressure Calibration and Pressure Post-Test tasks do not need to be performed, click **Finish** and then click **Abort Selected Tasks**.

7. Pressure Calibration:

Pressure Calibration

Step 1 Load a dry screened calibration set, then close the door.
Please pressurize the set to 0 psi on the patient side.

Enter Set Serial Number:

The set serial number is identified on the label attached to the 8100-PCS set. It is the number that follows the most recent date on the label. When entering the serial number, do not enter the decimal point.

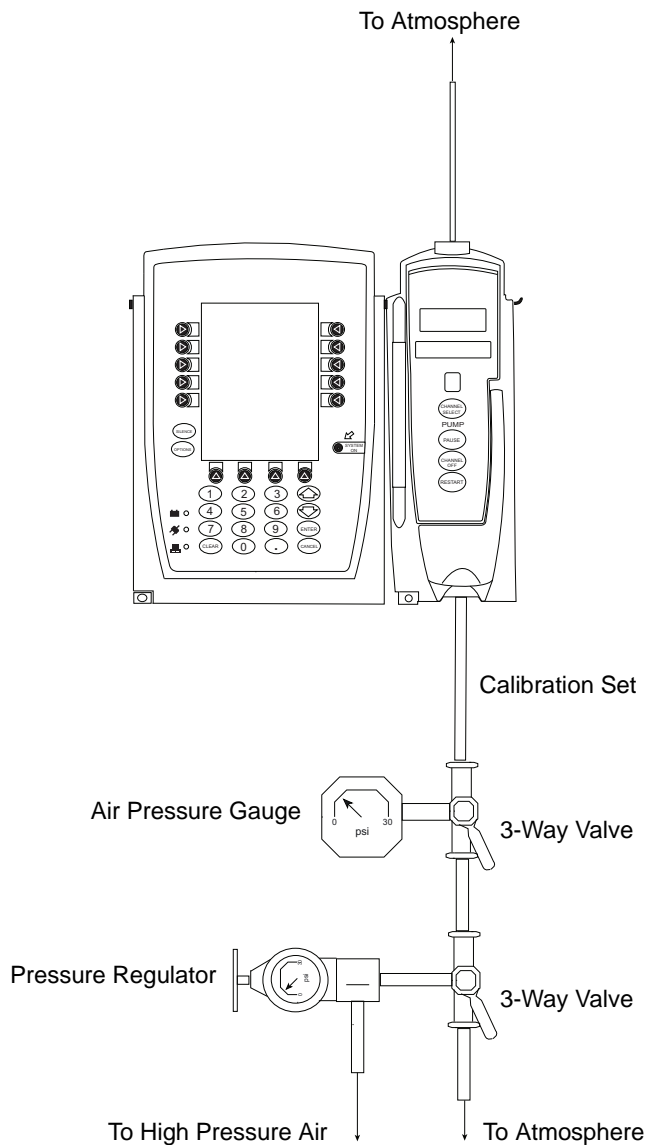
Patient-Side Occlusion Pressure Calibration Setup—Pump Module

Use the Model 8100-PCS pressure calibration set and the following setup to perform the patient-side occlusion pressure calibration (Task: Pressure Calibration).

Do not use the pressure gauge in peak hold mode during pressure calibration. Use the peak hold mode only during occlusion pressure tests.

CAUTION

- Pressure calibration sets are valid for 20 calibrations. Replace pressure calibration sets after 20 uses.
- To ensure accurate pressure calibration, use only pressure calibration sets.



Step 2 System Acclimation
System acclimation takes 30 seconds: 2

Step 3 Please enter the pressure reading from the gauge.
Acceptable range for the pressure is -0.1 to 0.1 psi.

Enter reading : psi

Step 4 Summary of the pressure parameters

Acceptable Upstream voltages:	1.835-2.830 volts
Actual Upstream voltage:	2.293 volts
Upstream sensor Passed	
Acceptable Downstream voltages:	1.803-2.674 volts
Actual Downstream voltage:	2.1526 volts
Downstream stream sensor Passed	
Pump Module Passed	

Step 5 Please pressurize the set to 10 psi on the patient side and then enter the pressure reading from the gauge.
Acceptable range for the pressure is 9.8 to 10.2.

Enter reading : psi

Step 6 Summary of the pressure parameters

Expected Downstream System Gain values:	0.1048-0.1464 volts/psi
Actual Downstream System Gain value:	0.1277 volts/psi
The Downstream System Gain is within the acceptable range.	

Step 7 Summary of results


Upstream pre-load voltage:	2.293 volts
Downstream pre-load voltage:	2.1526 volts
Downstream System Gain value:	0.1277 volts/psi
Calibration Complete.	

Step 8 A Pass result will be recorded for this device.

Release the pressure and remove the 8100-PCS set.

8. Pressure Post-Test:


Set up the Pump Module for the test (see *Patient-Side Occlusion Pressure Test Setup* on page 55).

 **Pressure Post-Test**

Step 1 Load a primed regular set and close the door. Ensure that the pressure gauge is set to zero.

Step 2 Wait until 0.4 ml has been infused before causing a Patient Side Occlusion.

VTBI	4 ml
Rate	125 ml/h
Elapsed Time	13 secs
Infused Volume	0.4 ml



After 0.4 ml has been delivered, simulate a patient-side occlusion by flipping the stop cock on the 3-way valve so that fluid is pumping toward the pressure transducer

Step 3 Pump Occluded. Enter the pressure reading at occlusion.

Pressure reading: psi

Enter the pressure reading at the time of Occlusion.
 Passing values are 7.7 to 12.7 psi.

Step 4 A Pass result will be recorded for this device.

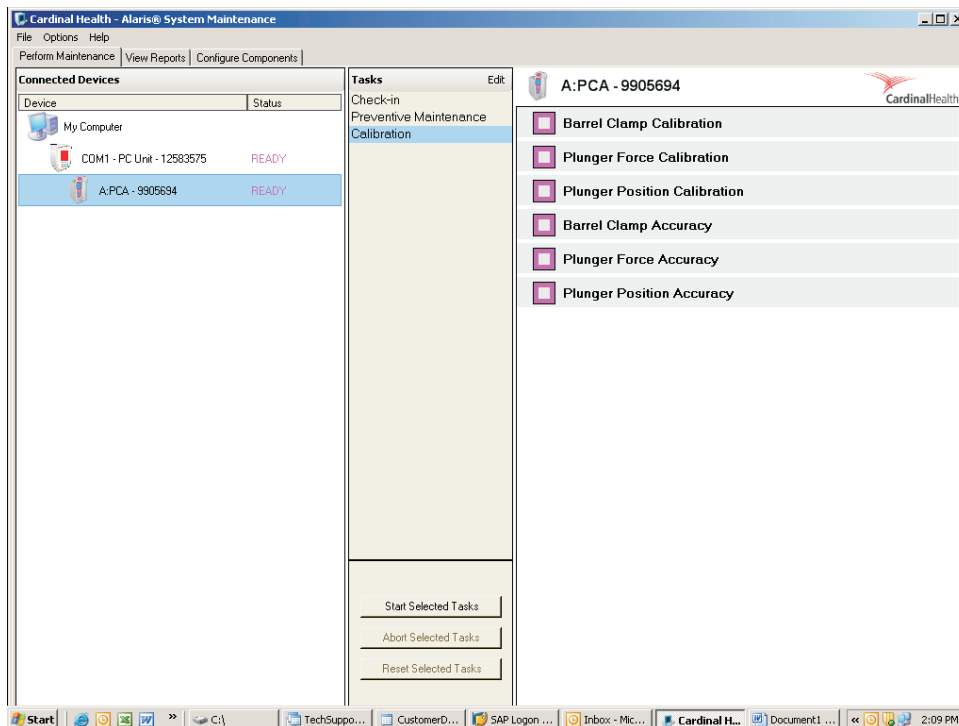
PCA Module Calibration

Test Equipment

Test Equipment	Manufacturer	Model/Part Number	Application
gauge, syringe height	CareFusion	148181-100	plunger position calibration / verification
kit, calibration tools ^a	CareFusion	10010692	all tools required for calibration and verification
kit, force sensor	CareFusion	10010691	force sensor calibration / verification
kit, syringe sizer and height gauge	CareFusion	148182-100	plunger position / barrel size calibration / verification
kit, syringe sizer gauge	CareFusion	148180-100	barrel size calibration / verification

a. To check the spring cartridge verification, preload the cartridge with 0.50 ± 0.05 lbs. Measure the deflection from the preload point after adding 19.5 lbs additional load (20 ± 0.05 lbs total). Acceptable deflection is within 0.891-0.940 inches. Use of Instron or equivalent is recommended.

1. Click **Perform Maintenance** tab.
2. In the Connected Devices pane, click the PCA Module to be tested.
3. Click **Calibration** in the Tasks list (double-click to immediately begin first task).



- Click the calibration task that needs to be performed and follow the instructions displayed on the screen.

NOTE

See *Syringe Module and PCA Module Tasks* on page 178 for an explanation of each task.

- Barrel Clamp Calibration:

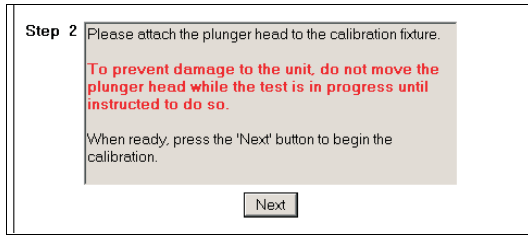
Insert the 10 mm diameter fixture behind the barrel clamp and close the barrel clamp on the fixture.

Remove the 10 mm diameter fixture, insert the 20 mm diameter fixture, and close the barrel clamp on the fixture. A popup alerts you if the correct size fixture is not installed.

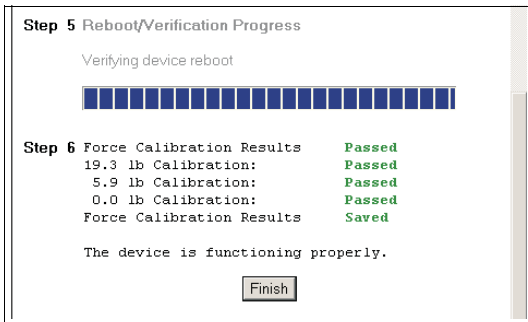
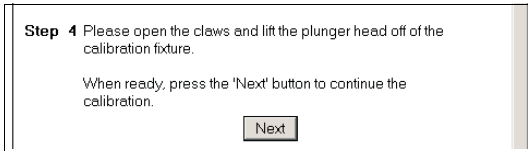
Remove the 20 mm diameter fixture, insert the 30 mm diameter fixture, and close the barrel clamp on the fixture. A popup alerts you if the correct size fixture is not installed.

- Plunger Force Calibration:

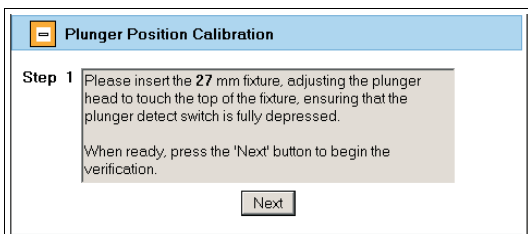
IMPORTANT: To avoid damage to the plunger header, ensure that the plunger header is not attached to the calibration fixture.



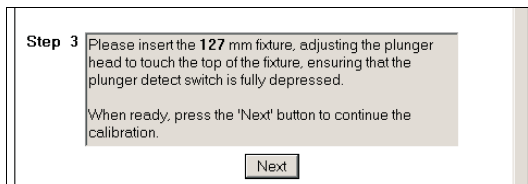
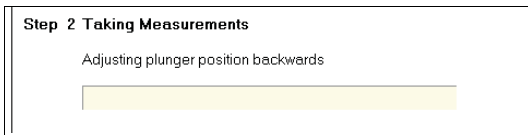
IMPORTANT: To avoid instrument damage, do not move the plunger head during this test.



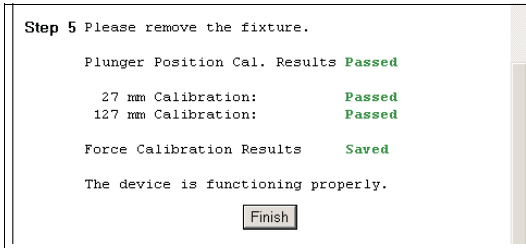
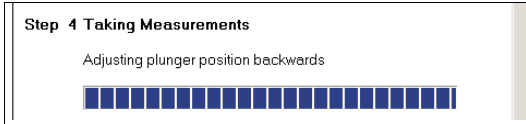
7. Plunger Position Calibration:



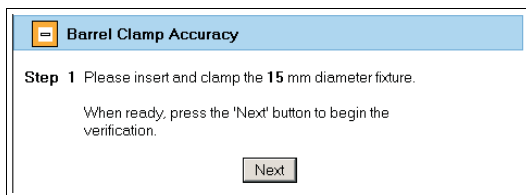
Follow the instructions on the screen and then close the barrel clamp on the 27 mm fixture before pressing **Next**.



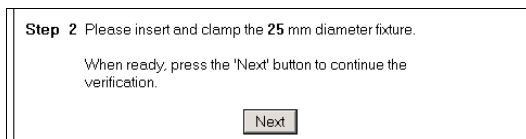
Remove the 27 mm diameter fixture, insert the 127 mm diameter fixture, close the barrel clamp on the fixture. A popup alerts you if the correct size fixture is not installed.



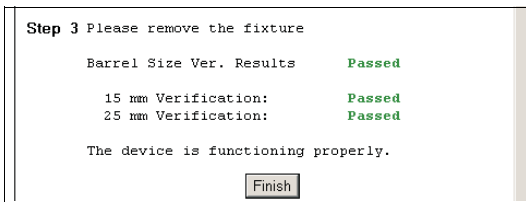
8. Barrel Clamp Accuracy:



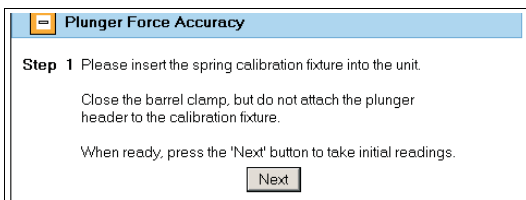
Insert the 15 mm diameter fixture behind the barrel clamp and close the barrel clamp on the fixture.



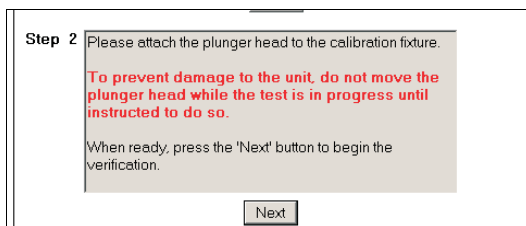
Remove the 15 mm diameter fixture, insert the 25 mm diameter fixture, and close the barrel clamp on the fixture. A popup alerts you if the correct size fixture is not installed.



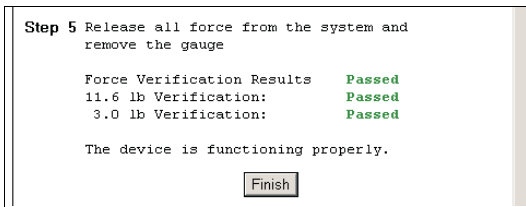
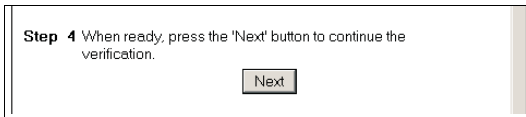
9. Plunger Force Accuracy:



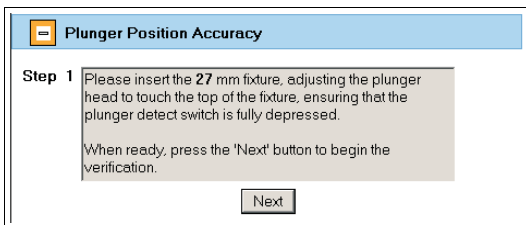
IMPORTANT: To avoid damage to the plunger header, ensure that the plunger header is not attached to the calibration fixture.



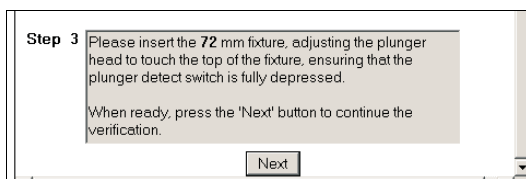
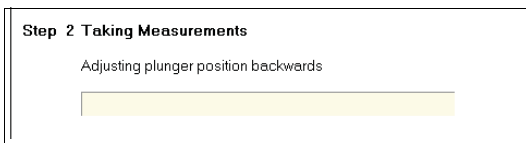
IMPORTANT: To avoid instrument damage, do not move the plunger head during this test.



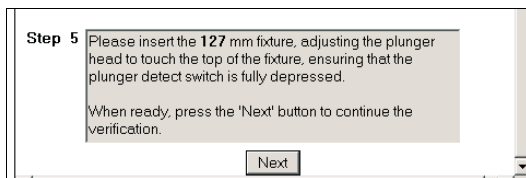
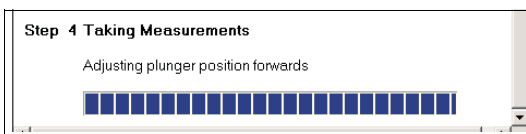
10. Plunger Position Accuracy:



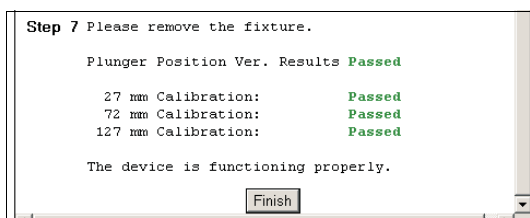
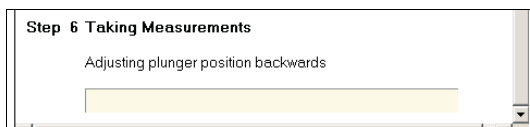
Follow the instructions on the screen and then close the barrel clamp on the 27 m fixture before pressing **Next**.



Remove the 27 mm diameter fixture, insert the 72 mm diameter fixture, close the barrel clamp on the fixture. A popup alerts you if the correct size fixture is not installed.



Remove the 72 mm diameter fixture, insert the 127 mm diameter fixture, close the barrel clamp on the fixture. A popup alerts you if the correct size fixture is not installed.



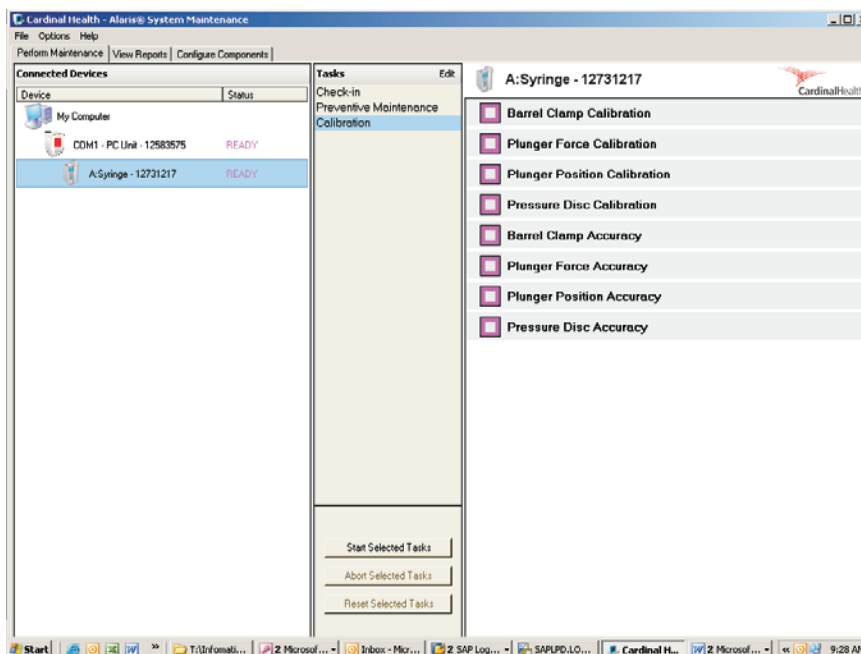
Syringe Module Calibration

Test Equipment

Test Equipment	Manufacturer	Model/Part Number	Application
gauge, syringe height	CareFusion	148181-100	plunger position calibration / verification
kit, calibration tools ^a	CareFusion	10010692	all tools required for calibration and verification
kit, force sensor	CareFusion	10010691	force sensor calibration / verification
kit, syringe sizer and height gauge	CareFusion	148182-100	plunger position / barrel size calibration / verification
kit, syringe sizer gauge	CareFusion	148180-100	barrel size calibration / verification
tubing, silicone	CareFusion	303109 or equivalent	pressure test
valve, 3-way	CareFusion	97555 or equivalent	pressure test

a. To check the spring cartridge verification, preload the cartridge with 0.50 ± 0.05 lbs. Measure the deflection from the preload point after adding 19.5 lbs additional load (20 ± 0.05 lbs total). Acceptable deflection is within 0.891-0.940 inches. Use of Instron or equivalent is recommended.

1. Click **Perform Maintenance** tab.
2. In the Connected Devices pane, click the Syringe Module to be tested.
3. Click **Calibration** in the Tasks list (double-click to immediately begin first task).



- Click the calibration task that needs to be performed and follow the instructions displayed on the screen.

NOTE

See *Syringe Module and PCA Module Tasks* on page 178 for an explanation of each task.

5. Barrel Clamp Calibration:

Barrel Clamp Calibration

Step 1 Please insert and clamp the **10 mm** diameter fixture.

When ready, press the 'Next' button to begin the calibration.

Step 2 Please insert and clamp the **20 mm** diameter fixture.

When ready, press the 'Next' button to continue the calibration.

Step 3 Please insert and clamp the **30 mm** diameter fixture.

When ready, press the 'Next' button to continue the calibration.

Step 4 Please remove the fixture and rotate the barrel clamp into the **Open Clamp** position.

When ready, press the 'Next' button to continue the calibration.

Step 5 Reboot/Verification Progress

Rebooting device

Step 6 Barrel Size Cal. Results **Passed**

10 mm Calibration: **Passed**

20 mm Calibration: **Passed**

30 mm Calibration: **Passed**

Calibration 1: **Saved**

Calibration Open Clamp: **Passed**

Calibration 2: **Saved**

The device is functioning properly.

6. Plunger Force Calibration:

Plunger Force Calibration

Step 1 Please insert the spring calibration fixture into the unit.

Close the barrel clamp, but do not attach the plunger header to the calibration fixture.

When ready, press the 'Next' button to take initial readings.

IMPORTANT: To avoid damage to the plunger header, ensure that the plunger header is not attached to the calibration fixture.

Step 2 Please attach the plunger head to the calibration fixture.


To prevent damage to the unit, do not move the plunger head while the test is in progress until instructed to do so.

When ready, press the 'Next' button to begin the calibration.

IMPORTANT: To avoid instrument damage, do not move the plunger head during this test.

Step 3 Calibration Progress

Adjusting to starting position




Step 4 Please open the claws and lift the plunger head off of the calibration fixture.

When ready, press the 'Next' button to continue the calibration.

Step 5 Reboot/Verification Progress

Verifying device reboot



Step 6 Force Calibration Results

19.3 lb Calibration:	Passed
5.9 lb Calibration:	Passed
0.0 lb Calibration:	Passed
Force Calibration Results	Saved

The device is functioning properly.

7. Plunger Position Calibration:

Plunger Position Calibration

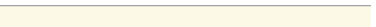
Step 1 Please insert the 27 mm fixture, adjusting the plunger head to touch the top of the fixture, ensuring that the plunger detect switch is fully depressed.

Do not use a pressure disc for this test.

When ready, press the 'Next' button to begin the verification.

Step 2 Taking Measurements

Adjusting plunger position backwards




Step 3 Please insert the 127 mm fixture, adjusting the plunger head to touch the top of the fixture, ensuring that the plunger detect switch is fully depressed.

When ready, press the 'Next' button to continue the calibration.

Step 4 Taking Measurements

Adjusting plunger position backwards



Step 5 Please remove the fixture.

Plunger Position Cal. Results **Passed**

27 mm Calibration: **Passed**
 127 mm Calibration: **Passed**

Force Calibration Results **Saved**

The device is functioning properly.

Finish

8. Pressure Disc Calibration:

Set up the Syringe Module for pressure calibration (see *Pressure Test/Calibration Setup* on page 75).

Pressure Disc Calibration

Step 1 Verify that the pressure gauge is set to mmHg. Install the test set in to the module and connect the set to the pressure gauge.

Apply pressure until the pressure gauge reads **1000** mmHg +/- 5 mmHg.

Next

Step 2 Reduce the pressure until the pressure gauge reads **400** mmHg +/- 5 mmHg.


Next

Step 3 Reduce the pressure until the pressure gauge reads **50** mmHg +/- 2 mmHg.

Next

Step 4 Reboot/Verification Progress

Verifying device reboot



Step 5 Pressure Calibration Results **Passed**

1000 mmHg Calibration: **Passed**
 400 mmHg Calibration: **Passed**
 50 mmHg Calibration: **Passed**
 Calibration: **Saved**

The device is functioning properly.

Finish

9. Barrel Clamp Accuracy:

Barrel Clamp Accuracy

Step 1 Please insert and clamp the 15 mm diameter fixture.
When ready, press the 'Next' button to begin the verification.

Step 2 Please insert and clamp the 25 mm diameter fixture.
When ready, press the 'Next' button to continue the verification.

Step 3 Please remove the fixture

Barrel Size Ver. Results	Passed
15 mm Verification:	Passed
25 mm Verification:	Passed

The device is functioning properly.

10. Plunger Force Accuracy:

Plunger Force Accuracy

Step 1 Please insert the spring calibration fixture into the unit.
Close the barrel clamp, but do not attach the plunger header to the calibration fixture.
When ready, press the 'Next' button to take initial readings.

Step 2 Please attach the plunger head to the calibration fixture.

To prevent damage to the unit, do not move the plunger head while the test is in progress until instructed to do so.

When ready, press the 'Next' button to begin the verification.

Step 3 Verification Progress

Adjusting to starting position

Step 4 When ready, press the 'Next' button to continue the verification.

Step 5 Release all force from the system and remove the gauge

Force Verification Results	Passed
11.6 lb Verification:	Passed
3.0 lb Verification:	Passed

The device is functioning properly.

IMPORTANT: To avoid damage to the plunger header, ensure that the plunger header is not attached to the calibration fixture.

IMPORTANT: To avoid instrument damage, do not move the plunger head during this test.

11. Plunger Position Accuracy:

Plunger Position Accuracy


Step 1 Please insert the 27 mm fixture, adjusting the plunger head to touch the top of the fixture, ensuring that the plunger detect switch is fully depressed.

Do not use a pressure disc for this test.

When ready, press the 'Next' button to begin the verification.

Step 2 Taking Measurements

Adjusting plunger position backwards



Step 3 Please insert the 72 mm fixture, adjusting the plunger head to touch the top of the fixture, ensuring that the plunger detect switch is fully depressed.

When ready, press the 'Next' button to continue the verification.

Step 4 Taking Measurements

Adjusting plunger position forwards

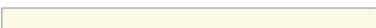


Step 5 Please insert the 127 mm fixture, adjusting the plunger head to touch the top of the fixture, ensuring that the plunger detect switch is fully depressed.

When ready, press the 'Next' button to continue the verification.

Step 6 Taking Measurements

Adjusting plunger position backwards



Step 7 Please remove the fixture.

Plunger Position Ver. Results **Passed**

27 mm Calibration: **Passed**

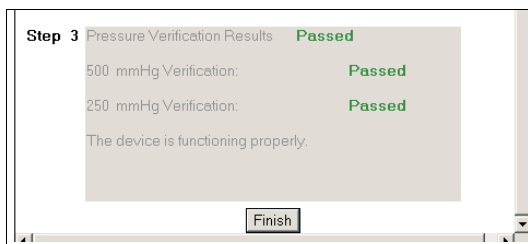
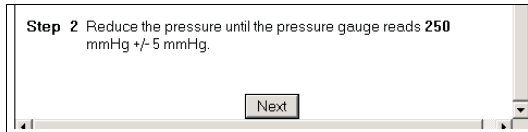
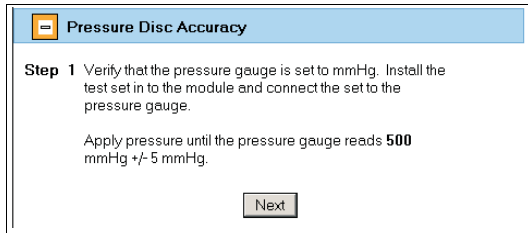
72 mm Calibration: **Passed**

127 mm Calibration: **Passed**

The device is functioning properly.

12. Pressure Disc Accuracy:

Set up the Syringe Module for the pressure test (see *Pressure Test/Calibration Setup* on page 75).

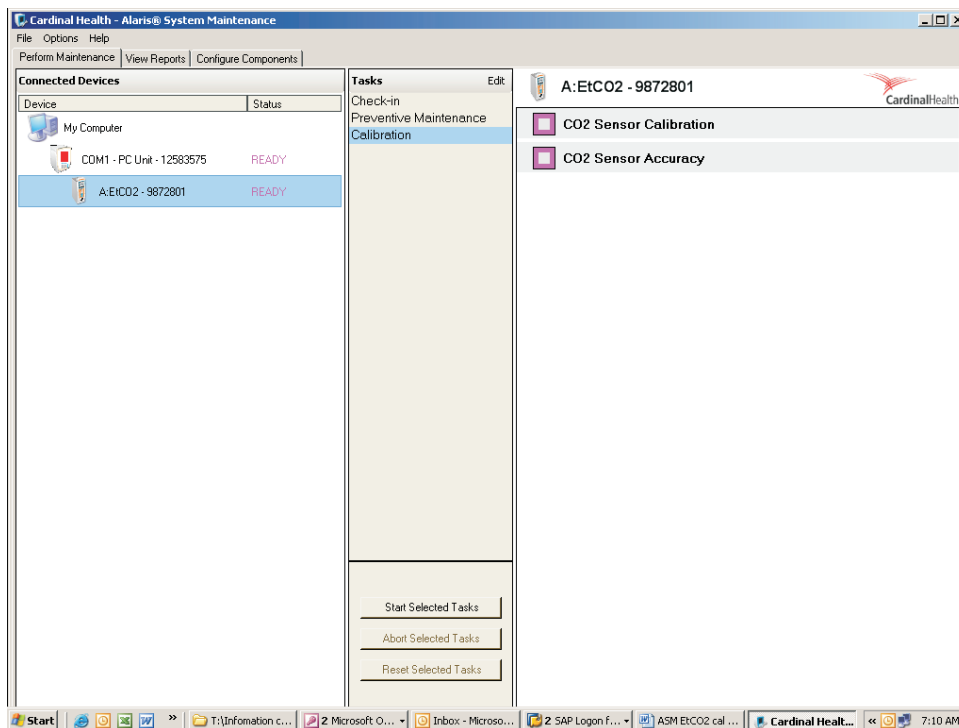


EtCO₂ Module Calibration

Test Equipment

Test Equipment	Manufacturer	Model Number	Application
EtCO ₂ pod (disposable)	Microstream®	007768	flow test/calibration
flow meter	Dwyer (www.dwyer-inst.com.au)	VFA-22, or equivalent	flow test/calibration
gas	Scott Medical	0304653SROBD, or equivalent gas with CO ₂ content of 4-6%	flow test/calibration

1. Click **Perform Maintenance** tab.
2. In the Connected Devices pane, click the EtCO₂ Module to be tested.
3. Click **Calibration** in the Tasks list (double-click to begin testing immediately).



4. Follow the instructions displayed on the screen for each test.

NOTE

See *EtCO₂ Module Tasks* on page 183 for an explanation of each task.

5. CO₂ Sensor Calibration:

Set up the EtCO₂ Module for the flow test/calibration (see *Flow Test/Calibration Setup* on page 89).

CO₂ Sensor Calibration

Step 1 Reminder: Do not calibrate this unit unless it has been warming up for at least 20 minutes.

Please connect a disposable and attach the gas source. When instructed, start the gas flowing and leave it flowing until instructed to stop.
 Caution: The gas flow rate must be above 500 mL/min. Use of a flow meter is recommended.

Next

Ensure that the module has been warming up for at least 20 minutes before performing calibration.

500 mL/min is equivalent to 0.5 liters per minute. Vent the flow meter to open air.

Step 2 Enter the percentage of CO₂ in the calibration gas (there should be 21% O₂ and the balance should be N₂).

% CO₂

Valid values are 4% to 6%

Next

Do not turn off the gas flow until instructed to in step 5.

Step 3 Start the gas flow

Next

Step 4 Running calibration. Keep the gas flowing.

Step 5 Calculating... Please turn off the gas source

Step 6 Test Result: Passed

Finish

6. CO₂ Sensor Accuracy:

CO₂ Sensor Accuracy

Step 1 Please connect a disposable and attach the gas source. When instructed, start the gas flowing and leave it flowing until instructed to stop.
 Caution: The gas flow rate must be above 500 mL/min. Use of a flow meter is recommended.

Next

500 mL/min is equivalent to 0.5 liters per minute. Vent the flow meter to open air.

Step 2 Enter the percentage of CO₂ in the verification gas (there should be 21% O₂ and the balance should be N₂).

%

Valid values are 4% to 6%

Next

Step 3 Start the gas flow

Next

Step 4 CO₂ sensor reading: 4.95

Step 5 Stop the gas and press 'Finish'

	Expected	Actual
CO ₂ Sensor value:	5.0%	4.95%
Acceptable range:	4.71% - 5.29%	
TEST PASSED		

Finish

Do not turn off the gas flow until instructed to in step 5.

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Chapter 11

Tasks

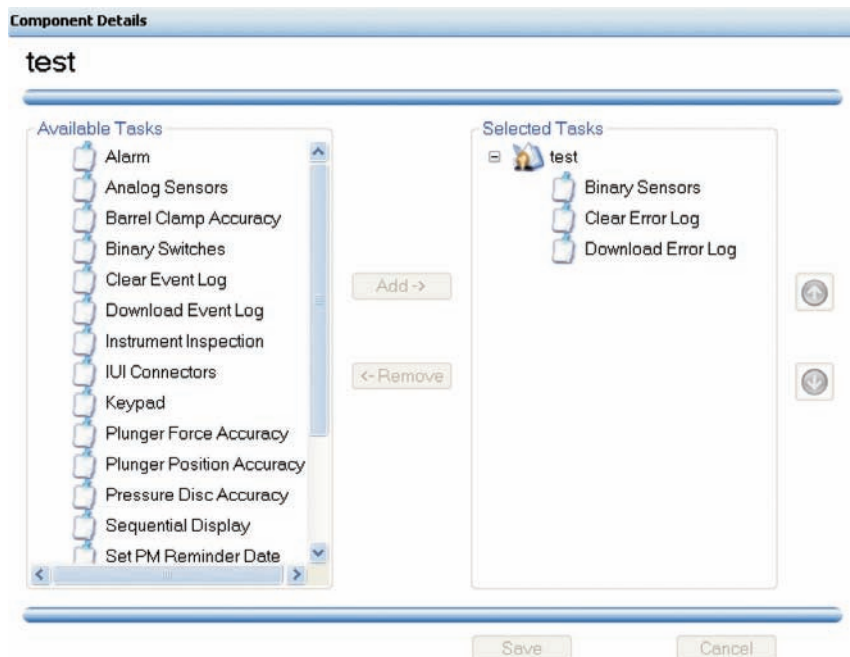
Task Groups

Create a New Task Group

Custom task groups can be defined and then added to the Task list. You cannot edit the standard task groups, which will always display in the Task list.

1. On the main window, click the **Configure Components** tab.
2. In the Components window, click the **Task Groups** tab.
3. At the top of the device type list, select a device type and click **New** (or right-click the device and click **New**).
4. Enter a name for the task group and press Enter.

The Component Details window is displayed.



5. If you wish to add a task, select the task from the **Available Tasks** list and click **Add** to add it to the **Selected Tasks** list for the device.
6. If you wish to remove a task, select the task in the **Selected Tasks** list and click **Remove** to remove it from the list.
7. Continue selecting tasks until you have collected all the needed tasks.
8. Use the up and down arrows to change the list order, if required.



9. Click **Save** to save changes.

The Task group name is added to the list of task groups in the Components window.

Delete a Task Group

Do *not* delete all task groups from the Tasks list. At least one task group must remain in the Tasks list.

1. On the main window, click the **Configure Components** tab.
2. In the **Components** window, click the **Task Groups** tab.
3. Right-click the name of a task group and click **Delete**.
4. Click **Yes** to confirm the deletion.

Rename a Task Group

1. On the main window, click the **Configure Components** tab.
2. In the Components window, right-click the name of a task group and click **Rename**.
3. Enter the new name in the dialog box and press Enter.

Tasks List

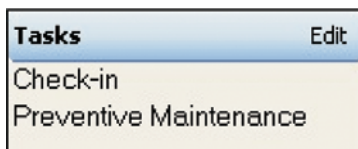
Edit Tasks List

You can edit the Tasks list to display tasks in addition to the standard task groups. The task list edit button must be enabled in Application Settings to be able to edit the tasks lists. You cannot edit the standard task groups, which always display in the Tasks list.

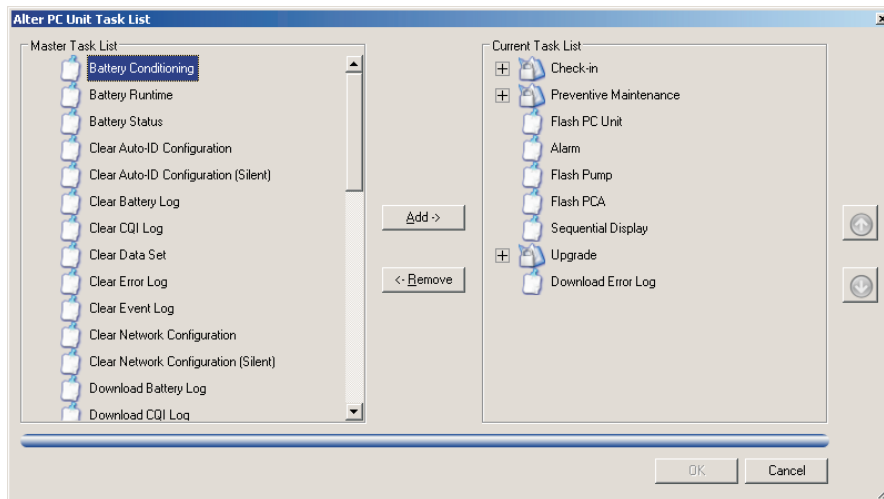
NOTE

For information on configuring task groups, see *Task Groups* on page 166.

1. On the main window, click the **Perform Maintenance** tab and select a device.
2. Click **Edit** in the Tasks list.



The Alter Task List dialog box is displayed for the selected device.



3. To add a task, select it in the Master Task List and click **Add** (or double-click it). To remove a task, select it in the Current Task List and click **Remove** (or double-click it).
4. Click **OK** to save the changes.

Change Tasks List Order

You can change the display order of tasks in a device Task list.

1. On the main window, click the **Perform Maintenance** tab and select a device.
2. Click **Edit** in the Tasks list.

The Alter Task List dialog box is displayed for the selected device.

3. Select a task in the Current Task List and use the arrows to move it up and down in the list.



4. Click **OK** to save changes.

Task Information

The following tables contain descriptions of tasks available for the PC Unit and any attached modules. The table also lists which standard task group each task is in.

PC Unit Tasks


PC Unit Tasks

Task	Standard Task Group	Description
Alarm	Check-in Preventive Maintenance	Tests audio and visual alarm prompts. You should be able to hear an audible tone from the PC Unit and see a flashing red light at the top of the attached module.
Battery Conditioning	(not in a standard task group)	Performs a sequence of battery drains and charges to condition the battery. This test can take eight hours or more to complete. It can be run overnight without being connected to the personal computer.
Battery Runtime	(not in a standard task group)	Tests the time a battery charge lasts (manually timed test). Nickel Metal Hydride (NiMH) batteries lose capacity over time (depending on usage factors, such as frequency and depth of discharge). Since the rate of loss increases as the capacity diminishes, CareFusion recommends replacing or reconditioning batteries if runtime for four channels is less than four hours (operating at 25 mL/h). Battery conditioning should be run to recondition the NiMH battery if the PC Unit has reduced battery capacity (for example, after repeated battery alarms).
Battery Status	(not in a standard task group)	Provides battery status information.
Clear Auto-ID Configuration	(not in a standard task group)	Clears Auto-ID Module configuration from PC Unit.
Clear Auto-ID Configuration (Silent)	(not in a standard task group)	Clears Auto-ID Module configuration (silent) from PC Unit.
Clear Battery Log	(not in a standard task group)	Clears battery log from PC Unit.
Clear CQI Log	(not in a standard task group)	Clears alerts stored in PC Unit.
Clear Data Set	(not in a standard task group)	Clears Data Set from PC Unit.
Clear Error Log	(not in a standard task group)	Clears error log.
Clear Event Log	(not in a standard task group)	Clears event log.
Clear Network Configuration	(not in a standard task group)	Clears network configuration from the PC Unit.
Clear Network Configuration (Silent)	(not in a standard task group)	Clears network configuration (silent) from the PC Unit.
Disable Wireless Configuration	(not in a standard task group)	Disables the wireless feature permanently on the PC Units. This is for PC Units that will be operating in a non-wireless environment.

PC Unit Tasks (Continued)

Task	Standard Task Group	Description
Download Battery Log	Upgrade	Downloads battery log from PC Unit. If logs are not downloaded and cleared regularly, storage space on the Alaris® System can be filled and older data will be overwritten as new events occur.
Download CQI Log	Upgrade	Downloads alerts stored in PC Unit to CQI database when wireless network is not available.
Download Error Log	Upgrade	Downloads error log that contains a log of abnormal device conditions due to hardware or software error. If you do not download and clear logs regularly, storage space on the Alaris® System can be filled and older data will be overwritten as new events occur.
Download Event Log	Upgrade	PC Unit and its modules create event logs, providing a history of key presses. These logs can be downloaded periodically to provide a database of module-specific events. If you do not download and clear logs regularly, storage space on the Alaris® System can be filled and older data will be overwritten as new events occur.
Flash Auto-ID	(not in a standard task group)	Updates module firmware from the Release Manifest File according to the overwrite setting.
Flash EtCO ₂	(not in a standard task group)	Updates module firmware from the Release Manifest File according to the overwrite setting.
Flash PC unit	Upgrade	Updates PC Unit firmware from the Release Manifest File according to the overwrite setting.
Flash PCA	(not in a standard task group)	Updates module firmware from the Release Manifest File according to the overwrite setting.
Flash Pump	(not in a standard task group)	Updates module firmware from the Release Manifest File according to the overwrite setting.
Flash SpO ₂	(not in a standard task group)	Updates module firmware from the Release Manifest File according to the overwrite setting.
Flash Syringe	(not in a standard task group)	Updates module firmware from the Release Manifest File according to the overwrite setting.
Ground Leakage	Check-in Preventive Maintenance	Tests that ground leakage is within accepted limits. See <i>Ground Leakage</i> : on page 45.
Ground Resistance	Check-in Preventive Maintenance	Tests that ground resistance is within accepted limits. See <i>Ground Resistance</i> : on page 45.
HiPot	(not in a standard task group)	Standard dielectric test of device insulation.
Instrument Inspection	Check-in Preventive Maintenance	Describes what to look for while examining a device. Description varies depending on module type selected.

PC Unit Tasks (Continued)

Task	Standard Task Group	Description
IUI Connectors	Check-in Preventive Maintenance	<p>Tests the proper functioning of IUI connectors that connect devices and PC Unit.</p> <p>You must connect a device to each connector of the PC Unit to test it. Connections can be tested one at a time or both at once.</p> 
Keypad	Check-in Preventive Maintenance	<p>Verifies that keys on the device keypad are functioning properly. Test instructions vary depending on module type selected.</p> <ul style="list-style-type: none"> Starting text for each key will be black. When a key press is registered, the text on the test for that key becomes bold and changes color to green. If a second key press is registered for that key, the text changes to red. If you see red text, rerun the test to verify the key is not stuck.
Network Connectivity		<p>Tests that the PC Unit can communicate wirelessly with the Alaris® network. You have the choice of testing the network parameters present on the PC Unit or temporary parameters, which runs test with the first parameter set configured in active configuration package.</p> <p>Use temporary parameters option to test a new parameter set before updating the PC Unit. The parameter set imported or created in current maintenance session is sent temporarily to the PC Unit. (After testing, clear parameters by power cycling unit or by rerunning the test.)</p> <p>This option runs only the first of the parameter sets to be tested. If you know that a particular set works with an Alaris® Server from the personal computer, make sure this parameter set appears first when defining network parameters.</p>
Nurse Call	Check-in Preventive Maintenance	<p>Verifies if a nurse call unit is installed. If one is installed, the test prompts you to press the Nurse Call button and verify that the nurse call is signaled.</p>
Power Supply	(not in a standard task group)	<p>Displays the results of the Power On Self Test (POST) from the power supply processor. During normal operation, if POST elements fail, the PC Unit does not power up. However, in Maintenance Mode, the device can power up to allow this test to isolate problems in the power supply processor.</p>
Sequential Display	(not in a standard task group)	<p>Tests the device display and status lights in sequence.</p>
Set PM Reminder Date	Check-in Preventive Maintenance	<p>Prompts you for a maintenance date and sets it for the selected device.</p>
Simultaneous Display	Check-in Preventive Maintenance	<p>Tests the device display and all status lights simultaneously.</p>
Transfer Auto-ID Configuration	(not in a standard task group)	<p>Transfers the Auto-ID Module configuration saved in the active configuration package to the PC Unit.</p>
Transfer Auto-ID Configuration (Silent)	Upgrade	<p>Transfers the Auto-ID Module configuration saved in the active configuration package to the PC Unit, in the Silent mode.</p>

PC Unit Tasks (Continued)

Task	Standard Task Group	Description
Transfer Data Set	Upgrade	<p>Transfers a Data Set to a PC Unit when it is not connected to the wireless network.</p> <p>Draft Data Sets and Approved Data Sets can be transferred for training or clinical interface testing purposes. When a Draft Data Set or an Approved Data Set is loaded, the message "This Device is Not for Human Use" is displayed on the PC Unit screen.</p> <p>Released Data Sets are only used after they are approved by the facility and are ready for human use.</p> <p>WARNING Unreleased Data Sets are transferable only for user interface testing and/or training and are not for use on patients.</p> <p>CAUTION</p> <ul style="list-style-type: none"> • If you have not yet downloaded the alerts from the Alaris® System, download them before you transfer the Data Set. • The new Data Set will not become active until you power-cycle the PC Unit and select New Patient.
Transfer Network Configuration	(not in a standard task group) For error handling, see <i>Transfer Network Configuration—Error Handling</i> on page 174.	Transfers the network configuration (parameter sets and Alaris® Server information) that are saved in the active configuration package to the PC Unit. Allows you to view, clear, and export the network configuration settings currently on the PC Unit.
Transfer Network Configuration (Silent)	Upgrade For error handling, see <i>Transfer Network Configuration—Error Handling</i> on page 174.	Transfers the network configuration (parameter sets and Alaris® Server information) to the PC Unit in a single step. The additional features that are available when using the Transfer Network Configuration task cannot be accessed when using this task.
Voltage Display	(not in a standard task group)	Shows the power supply and system voltage information and identifies values outside the acceptable range.

Transfer Network Configuration—Error Handling

PC Unit Main Processor & Wireless Main Version	Wireless RF Card	802.11d	802.11a	WEP	WPA PSK, TLS, TTLS, PEAP	WPA2 PSK, TLS, TTLS, PEAP	PSK		Encryption		Compatibility	
							ASCII	Hex	TKIP	AES	Partially Incompatible	Completely Incompatible
v9.1.x	Any supported brand of RF card or no RF card	N ¹	N ¹	Y	Y	N ¹	Y	N ¹	Y	N ¹	¹ Error popup: PC Unit embedded Wireless Main software version v9.1.X doesn't support profile: Profile X. Network Configuration will not be transferred to the connected PC Unit.	
v9.5.x	Symbol b	N ²	N ²	Y	Y	N ²	Y	Y	Y	N ²	² Information popup: Settings for Profile X are not supported for this particular device RF card.	² Error popup: RF Card on connected PC Unit is not supported or is not compatible with Network Configuration Provided. Network Configuration will not be transferred to the connected PC Unit.
	AmbiCom b/g	N ²	N ²	Y	Y	Y	Y	Y	Y	Y		
	Motorola a/b/g	Y	Y	Y	Y	Y	Y	Y	Y	Y	Y	
	No RF card	n/a ⁰	n/a ⁰	n/a ⁰	n/a ⁰	n/a ⁰	n/a ⁰	n/a ⁰	n/a ⁰	n/a ⁰	⁰ Error popup: No RF card was detected on this PC Unit. Network Configuration will not be transferred to the connected PC Unit.	

- **N**—No (not compatible).
- **Y**—Yes (compatible).
- **X**—Represents the software version or profile number that appears in the popup message.
- **n/a**—Not Applicable.
- **Partially Incompatible**—At least one of the network profiles is supported. When you click the **OK** button, the supported network settings are transferred to the PC Unit.
- **Completely Incompatible**—The network profile is not supported. The network settings will not be transferred to the PC Unit.

Pump Module Tasks

Unless otherwise specified, a disposable set must be loaded and the Pump Module door must be closed to run any Pump Module test. For instructions, see *Load Administration Set* on page 51.

Pump Module Tasks

Task	Standard Task Group	Description
Air-in-Line Sensor	Check-in Preventive Maintenance	Verifies Air-in-line detection and also that a dry set is detected.
Air-in-Line Sensor (Wet and Dry)	(not in a standard task group)	Air-in-line detection is verified using a wet set to detect sensing of wet or dry.
Alarm	Check-in Preventive Maintenance	See <i>Alarm</i> on page 170.
Analog Sensor	(not in a standard task group)	Displays readings from sensors that transmit analog measurements in real-time. Shows analog-to-digital readings of sensors with variable outputs. This view includes display of voltage outputs from pressure sensor strain gauges.
Binary Sensors	(not in a standard task group)	Shows 1 or 0 states of sensors that can be either ON or OFF.
Clear Error Log	(not in a standard task group)	See <i>Clear Error Log</i> on page 170.
Clear Event Log	(not in a standard task group)	See <i>Clear Event Log</i> on page 170.
Door Ajar-Flo-Stop Sensor	Check-in Preventive Maintenance	Tests sensor to verify that a door ajar and Flo-Stop conditions are detected.
Download Error Log	(not in a standard task group)	See <i>Download Error Log</i> on page 171.
Download Event Log	(not in a standard task group)	See <i>Download Event Log</i> on page 171.
Fluid-Side Occlusion	Check-in Preventive Maintenance	Tests that an occlusion in the fluid-side of the installed set is detected. For a diagram of the test setup, see <i>Fluid-Side Occlusion Test Setup</i> on page 57. CAUTION If the Pump Module fails the fluid-side occlusion test, it must be repaired.
Infuse Without Alarms	(not in a standard task group)	Runs an infusion at a user-defined rate and VTBI without activating alarms. During this test you can view the pumping mechanism with the door open to observe operation. While the infusion is running, open the door to examine infusion components and remove the set fitments or entire IV set.
Instrument Inspection	Check-in Preventive Maintenance	See <i>Instrument Inspection</i> on page 171.
IUI Connectors	Check-in Preventive Maintenance	See <i>IUI Connectors</i> on page 172.
Keypad	Check-in Preventive Maintenance	See <i>Keypad</i> on page 172.
Maximum Pressure	(not in a standard task group)	Determines the Pump Module's maximum mechanical pumping pressure by running an infusion to occlusion without activating a pressure alarm. Measured peak pressure is stored in the database.

Pump Module Tasks (Continued)

Task	Standard Task Group	Description
Patient-Side Occlusion	Check-in Preventive Maintenance	Tests that an occlusion in the patient side of the installed set is detected at a pressure that falls within calibration limits. For information about how to run the patient-side occlusion pressure test (including a diagram of the test setup), see <i>Patient-Side Occlusion Pressure Test Setup</i> on page 55. CAUTION If the Pump Module fails the fluid-side occlusion test, it must be repaired.
Power On Self Test	(not in a standard task group)	Invokes a power-on self test.
Pressure Calibration	Calibration	Pressure calibration is only required if the Pump Module fails the pressure calibration pre-test or after you replace the Pump Module. For information about running the pressure calibration test (including a diagram of the test setup), see <i>Patient-Side Occlusion Pressure Calibration Setup—Pump Module</i> on page 145. CAUTION <ul style="list-style-type: none"> To ensure accurate pressure calibration, use only pressure calibration sets. Pressure calibration sets are valid for 20 calibrations. Replace pressure calibration sets after 20 uses.
Pressure Post-Test	Calibration	Verifies that the calibration was successful.
Pressure Pre-Test	Calibration	Verifies that calibration is required.
Rate Accuracy	Check-in Preventive Maintenance	Checks that the accuracy of a module's rate of infusion falls within calibration limits. For more information about the rate accuracy test, including a diagram of the test setup, see <i>Rate Accuracy Test Setup</i> on page 52. Use the setup shown to perform the rate accuracy verification. Verify that the scale is calibrated according to the manufacturer's instructions. Prime the fluid lines and keep them free of air bubbles while running the test. CAUTION <ul style="list-style-type: none"> If the Pump Module fails the rate accuracy verification test, it must be calibrated. Failure to perform calibration tests might result in improper operation. Use distilled water at room temperature of 41°F to 104°F (5°C to 40°C). If water temperature is not within this temperature range, the readings might be inaccurate.
Rate Accuracy Post-Test	Calibration	Verifies that the calibration was successful.
Rate Accuracy Pre-Test	Calibration	Verifies that calibration is required.

Pump Module Tasks (Continued)

Task	Standard Task Group	Description
Rate Calibration	Calibration	<p>Rate accuracy calibration is only required if the Pump Module fails the rate accuracy calibration pre-test or after the pump component is replaced.</p> <p>Test the primes line to prevent an air-in-line warning from interrupting the test. For accurate results, ensure that the air is cleared from the line.</p> <p>For more information about running rate accuracy calibration tasks, including a diagram of the test setup, see <i>Rate Accuracy Test Setup</i> on page 52.</p> <p>CAUTION</p> <ul style="list-style-type: none"> To ensure accurate rate calibration, use only a rate calibration set. Use distilled water at room temperature of 41°F to 104°F (5°C to 40°C). If water temperature is not within this temperature range, the readings might be inaccurate. A rate accuracy calibration set is valid for 60 calibrations. After 60 uses, replace the set.
Sequential Display	(not in a standard task group)	See <i>Sequential Display</i> on page 172.
Set Loaded	(not in a standard task group)	Tests the sensor to verify proper set loading detected.
Set PM Reminder Date	Check-in Preventive Maintenance	See <i>Set PM Reminder Date</i> on page 172.
Simultaneous Display	Check-in Preventive Maintenance	See <i>Simultaneous Display</i> on page 172.
Time to Occlusion	(not in a standard task group)	Tests occlusions are detected within specified time limit.

Syringe Module and PCA Module Tasks

Syringe Module and PCA Module Tasks

Task	Standard Task Group	Description
Alarm	Check-in Preventive Maintenance	See <i>Alarm</i> on page 170.
Analog Sensors	(not in a standard task group)	<p>Enables the display readings of four sensors that transmit analog measurements from sensors in real-time.</p> <p>During the analog sensor test, you can move the head up and down to see height readings from the plunger position sensor, and open and close the clamp to see readings from the barrel size sensor. This test also loads a syringe, runs an infusion, and displays readings from the pressure and force sensors.</p> <p>This test displays sensor readings results:</p> <ul style="list-style-type: none"> • Pressure Sensor displays pressure readings in millimeters of mercury (mmHg). • Force Sensor displays current force applied to drive head in pounds (lbs). • Barrel Size Sensor displays outer diameter measurement of a loaded syringe in millimeters (mm). • Plunger Position Sensor displays syringe plunger height in millimeters (mm). <p>An occlusion warning alarm can occur if the selected rate is too high for the syringe and set. Larger diameter tubing offers less resistance and is therefore less likely to occlude during this test.</p>
Barrel Clamp Calibration	Calibration	<p>Requires three metal fixtures (10, 20, and 30 mm wide). Values returned by the calibration are flashed into the barrel size sensor.</p> <p>If the calibration fails, you are prompted to return the Syringe Module for repair.</p>
Barrel Clamp Accuracy	Calibration Check-in Preventive Maintenance	<p>This verification test uses a series of fixtures of a known diameter. Insert the fixtures according to on-screen instructions. Ensure that the fixtures seat vertically upright under the clamp and that the clamp rests snugly against the side of the fixture.</p> <p>Failing a verification test does not affect the calibration status of a Syringe Module or a PCA Module. To determine the calibration status, run the calibration settings test.</p>

Syringe Module and PCA Module Tasks (Continued)

Task	Standard Task Group	Description
Binary Sensors	(not in a standard task group)	<p>Displays the readings from six module sensors in real-time. During the binary sensors test you can insert, remove, push, pull, open, and close the working parts of the syringe and see in real-time what the Syringe Module detects.</p> <p>This test displays sensor readings results:</p> <ul style="list-style-type: none"> • Pressure Disc indicates the presence or absence of a pressure disc (Syringe Module only). • Flange indicates whether the syringe flange is detected. • Knob indicates whether the syringe plunger grippers (claws) are open or closed. • Split Nut indicates if the device is ready to infuse (open or closed). • Plunger indicates whether the drive head is in contact with a loaded syringe. • Clamp indicates whether the barrel clamp is open or closed.
Binary Switches	Check-in Preventive Maintenance	Displays readings from six module switches in real-time. During the binary switches test you can insert, remove, push, pull, open, and close the working parts of the syringe and see in real-time what the Syringe Module detects. Switches are the same as listed for the Binary Sensors test.
Clear Error Log	(not in a standard task group)	See <i>Clear Error Log</i> on page 170.
Clear Event Log	(not in a standard task group)	See <i>Clear Event Log</i> on page 170.
Download Error Log	(not in a standard task group)	See <i>Download Error Log</i> on page 171.
Download Event Log	(not in a standard task group)	See <i>Download Event Log</i> on page 171.
Instrument Inspection	Check-in Preventive Maintenance	See <i>Instrument Inspection</i> on page 171.
IUI Connectors	Check-in Preventive Maintenance	See <i>IUI Connectors</i> on page 172.
Keypad	Check-in Preventive Maintenance	See <i>Keypad</i> on page 172.

Syringe Module and PCA Module Tasks (Continued)

Task	Standard Task Group	Description
Plunger Force Accuracy	Calibration Check-in Preventive Maintenance	Tests force the sensor to verify that the module detects an occlusion alarm accurately. To generate force, the test moves drive head down against a spring calibration fixture. CAUTION <ul style="list-style-type: none"> To avoid damage to the module, ensure that the plunger grippers (claws) surround the top of the spring calibration fixture when instructed to attach them. To avoid damage to the module and possible injury, do not open the grippers or attempt to raise the drive head while the test is running. Open the grippers only when prompted by the software.
Plunger Force Calibration	Calibration	Measures force readings at 3 points from 0 to about 20 lbs of force. Values returned by three-point calibration are flashed into the syringe force sensor. If the calibration fails, you are prompted to return the Syringe Module to the service depot for repair. CAUTION To avoid damage to the module and possible injury, do not open the grippers or attempt to raise the drive head while the test is running. Open the grippers only when prompted by the software.
Plunger Position Accuracy	Calibration Check-in Preventive Maintenance	Uses fixtures to verify that the system correctly recognizes the plunger position by seating it on fixtures of known heights.
Plunger Position Calibration	Calibration	Requires two metal fixtures (27-mm and 127-mm lengths). When prompted, load each fixture as a syringe. The test moves the drive head down onto the top of each fixture to a preload that ensures firm contact has been made. The sensor is adjusted for the actual height of each fixture.
Pressure Disc Accuracy Verification (Syringe Module only)	Calibration Check-in Preventive Maintenance	Determines whether the Syringe Module pressure sensor requires calibration. It uses a digital pressure gauge and a pressure source to generate a known pressure in the Syringe Module, and then reads the pressure sensor at verification point(s).
Pressure Disc Calibration (Syringe Module only)	Calibration	Calibrates the Syringe Module pressure sensor. It uses a digital pressure gauge and a pressure source to generate a known pressure in the Syringe Module, and then reads the pressure sensor at the calibration points.

Syringe Module and PCA Module Tasks (Continued)

Task	Standard Task Group	Description
Sequential Display	(not in a standard task group)	See <i>Sequential Display</i> on page 172.
Set PM Reminder Date	Check-in Preventive Maintenance	See <i>Set PM Reminder Date</i> on page 172.
Simultaneous Display	Check-in Preventive Maintenance	See <i>Simultaneous Display</i> on page 172.

SpO₂ Module Tasks

SpO₂ Module Tasks

Task	Standard Task Group	Description
Alarm	Check-in Preventive Maintenance	See <i>Alarm</i> on page 170.
Clear Error Log	(not in a standard task group)	See <i>Clear Error Log</i> on page 170.
Clear Event Log	(not in a standard task group)	See <i>Clear Event Log</i> on page 170.
Download Error Log	(not in a standard task group)	See <i>Download Error Log</i> on page 171.
Download Event Log	(not in a standard task group)	See <i>Download Event Log</i> on page 171.
Instrument Inspection	Check-in Preventive Maintenance	See <i>Instrument Inspection</i> on page 171.
IUI Connectors	Check-in Preventive Maintenance	See <i>IUI Connectors</i> on page 172.
Keypad	Check-in Preventive Maintenance	See <i>Keypad</i> on page 172.
Patient Cable Alarm	Check-in Preventive Maintenance	Tests that the Patient Cable Alarm is functioning. Attach the patient cable and remove it when the test tells you to do so.
Patient Lead Electrical Leakage	Check-in Preventive Maintenance	Test for electrical leakage in patient lead.
Power On Self Test	Check-in Preventive Maintenance	Invokes a power on self test.
Pulse Rate/Saturation	Check-in Preventive Maintenance	Simulates known pulse rates and blood oxidation levels using the SpO ₂ Module finger simulator.
Sequential Display	(not in a standard task group)	See <i>Sequential Display</i> on page 172.
Set PM Reminder Date	Check-in Preventive Maintenance	See <i>Set PM Reminder Date</i> on page 172.
Simultaneous Display	Check-in Preventive Maintenance	See <i>Simultaneous Display</i> on page 172.
Speaker	Check-in Preventive Maintenance	Causes the speaker on the SpO ₂ Module to beep.

EtCO₂ Module Tasks

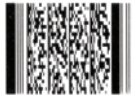
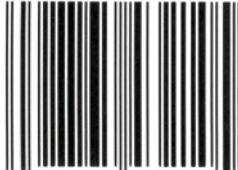
EtCO₂ Module Tasks

Task	Standard Task Group	Description
Alarm	Check-in Preventive Maintenance	See <i>Alarm</i> on page 170.
Ambient Pressure Sensor	(not in a standard task group)	Compares the ambient pressure sensor reading with an outside source. Requires an accurate source of barometric pressure.
Clear Error Log	(not in a standard task group)	See <i>Clear Error Log</i> on page 170.
Clear Event Log	(not in a standard task group)	See <i>Clear Event Log</i> on page 170.
CO ₂ Sensor Accuracy	Calibration Check-in Preventive Maintenance	To run this test, enter the percentage of CO ₂ gas in verification gas supply between 4% and 6%. Verify that gas flow is greater than 0.5 liters/minute. CAUTION The flow rate out of the gas source and into the EtCO ₂ Module must be greater than 0.5 liters/minute to ensure accurate readings. It is recommended that you attach a flow meter to the breather line.
CO ₂ Sensor Calibration	Calibration Preventive Maintenance	A verifiable gas source is required to run this calibration. CAUTION The flow rate out of the gas source and into the EtCO ₂ Module must be greater than 0.5 liters/minute to ensure accurate readings. It is recommended that you attach a flow meter to the breather line.
Disposable Connected	(not in a standard task group)	Verifies that the system can detect a disposable connection.
Download Error Log	(not in a standard task group)	See <i>Download Error Log</i> on page 171.
Download Event Log	(not in a standard task group)	See <i>Download Event Log</i> on page 171.
Instrument Inspection	Check-in Preventive Maintenance	See <i>Instrument Inspection</i> on page 171.
IUI Connectors	Check-in Preventive Maintenance	See <i>IUI Connectors</i> on page 172.
Keypad	Check-in Preventive Maintenance	See <i>Keypad</i> on page 172.
Leak Down	Check-in Preventive Maintenance	Tests for leakage from the gas pathway. For a diagram of the test setup, see <i>Leak-Down Test</i> on page 87.

EtCO₂ Module Tasks (Continued)

Task	Standard Task Group	Description
Sequential Display	(not in a standard task group)	See <i>Sequential Display</i> on page 172.
Set PM Reminder Date	Check-in Preventive Maintenance	See <i>Set PM Reminder Date</i> on page 172.
Simultaneous Display	Check-in Preventive Maintenance	See <i>Simultaneous Display</i> on page 172.

Auto-ID Module and Handheld Scanner Tasks**Auto-ID Module and Handheld Scanner Tasks**

Task	Standard Task Group	Description
2-D Scan	Check-in Preventive Maintenance	
Instrument Inspection	Check-in Preventive Maintenance	See <i>Instrument Inspection</i> on page 171.
IUI Connectors	Check-in Preventive Maintenance	See <i>IUI Connectors</i> on page 172.
Keypad	Check-in Preventive Maintenance	See <i>Keypad</i> on page 172.
LED	Check-in Preventive Maintenance	Verifies that LEDs on the module are functioning properly.
Linear Scan	(not in a standard task group)	
Scanner Connector	(not in a standard task group)	Checks the handheld scanner connector on the Auto-ID Module.
Scanner Functionality	Check-in Preventive Maintenance	Verifies proper scanner operation by performing a number of scans under controlled conditions. A minimum number of scans must be successful in order for a test to pass.
Set PM Reminder Date	Check-in Preventive Maintenance	See <i>Set PM Reminder Date</i> on page 172.

Test Equipment

Test Equipment	Manufacturer	Model/Part Number	Application
analyzer, electrical safety	Fluke BioMedical	232D or equivalent	PC Unit: AC wiring / instrument grounding and leakage tests. SpO ₂ patient lead electrical leakage test.
cable, Masimo® Simulator	Clinical Dynamics	SS-MAS	SpO ₂ pulse rate/saturation
cable, Nellcor® Oximax® Simulator	Clinical Dynamics	SS-NEL	SpO ₂ pulse rate/saturation
door key	CareFusion	10010964	PCA Module: lock and unlock PCA Module door
Dose Request Cord	CareFusion	10013795	PCA Module display test
EtCO ₂ pod (disposable)	Microstream®	007768	EtCO ₂ flow test/calibration
extension set, with pressure disc, small bore	CareFusion	10014917	Syringe Module pressure test
flow meter	Dwyer (www.dwyer-inst.com.au)	VFA-22, or equivalent	EtCO ₂ flow test/calibration
gas	Scott Medical	0304653SROBD, or equivalent gas with CO ₂ content of 4-6%	EtCO ₂ flow test/calibration
gauge, pressure, digital (peak hold)	Either of the following: <ul style="list-style-type: none"> Heise (www.heise.com) Ashcroft (www.ashcroft.com) 	<ul style="list-style-type: none"> PTE-1 (accuracy from 0.1 to 0.025% span) 2089, 2086, or 2084 (accuracy from ±0.05%, 0.10%, or 0.25% of span) or an equivalent gauge with: (a) unit of measure in mmHg (b) accuracy of ±1% (c) range of 0-1500 mmHg 	Pump Module pressure test and Syringe Module pressure test / calibration
gauge, syringe height	CareFusion	148181-100	PCA Module and Syringe Module plunger position calibration / verification
hemostat	N/A	N/A	Pump Module fluid-side occlusion test
IV pole, standard	CareFusion	903-0336 or equivalent	Pump Module rate, pressure, and occlusion tests
IV sets <ul style="list-style-type: none"> IV set, standard, without check valve IV set, calibration IV set, calibration 	<ul style="list-style-type: none"> CareFusion CareFusion CareFusion 	<ul style="list-style-type: none"> 2210-0500 8100-RCS 8100-PCS 	Pump Module rate, pressure, and occlusion tests
IV solution container (bag preferred)	N/A	N/A	Pump Module rate and pressure tests

Test Equipment	Manufacturer	Model/Part Number	Application
kit, calibration tools	CareFusion	10010692	PCA Module and Syringe Module: all tools required for calibration and verification
kit, force sensor	CareFusion	10010691	PCA Module and Syringe Module: force sensor calibration / verification
kit, syringe sizer and height gauge	CareFusion	148182-100	PCA Module and Syringe Module: plunger position / barrel size calibration / verification
kit, syringe sizer gauge	CareFusion	148180-100	PCA Module and Syringe Module: barrel size calibration / verification
luer lock, female, 1/8"-27 NPT	Cole-Parmer (www.coleparmer.com)	K-45503-78 or equivalent	Pump Module pressure test
pressure transducer simulator/tester	Utah Medical	Delta-Cal 650-95,0 or equivalent	EtCO ₂ leak-down test
reducer, female, 1/4" NPT (F) x 1/8" NPT (F)	Cole-Parmer (www.coleparmer.com)	K-06349-91 or equivalent	Pump Module pressure test
scale, digital	Acculab (www.accumlab.com)	VIC-212 (VICON Series) or equivalent with accuracy of $\pm 0.01g$	Pump Module rate and pressure tests
SpO ₂ simulator	Clinical Dynamics	SmartSat Pulse Oximetry Analyzer SS-100A, or equivalent	SpO ₂ pulse rate/saturation test
T-fitting	CareFusion	303815 or equivalent	Pump Module pressure test
tubing, silicone	CareFusion	303109 or equivalent	Pump Module, Syringe Module, and EtCO ₂ Module pressure test
valve, 3-way	CareFusion	97555 or equivalent	Pump Module, Syringe Module, and EtCO ₂ Module pressure test

Chapter 12

Reports and Logs

Reports

This section describes the report types and provides a procedure on how to run the reports.

Report Types

A number of maintenance reports are available from data stored in the System Maintenance Software database and/or the Maintenance Log database.

Activity	Reports on all activity performed on selected devices including tests, firmware updates, Data Set transfers, and Alert log downloads.
Asset	Reports on all device types in the database, showing serial number, PM due date, last calibration date, when firmware was last updated, and version numbers of installed firmware.
Log—Battery	Reports internal and external events relating to the battery, such as charging state and other battery status conditions.
Log—Error	Reports on abnormal events, such as internal software and hardware malfunctions.
Log—Event	Reports all operating events, such as keyboard presses, alarms, module attachments, external communications, plus infusion starts and stops. It might also contain additional information in coded form about changes that occurred on the display screen and other internal software events.
PM Due	Charts when selected module types are due for preventive maintenance.
Test—All	Reports on all tests performed on selected devices.
Test—Check-in	Reports on check-in activity on selected devices.
Test—PM	Reports on preventive maintenance activity on selected devices.

Run Reports

To run reports from the System Maintenance Software database:

1. On the main window, click the **View Reports** tab.
2. Select report type in the **Reports** pane.
3. Select devices by clicking the **Select Devices** button. In the Device Selection dialog box, you can use the following techniques:
 - Ctrl-click multiple devices to select them.
 - Shift-click a range of devices to select them.
 - Click **Filter** to select specific model numbers or PM Due dates for display.
 - Click **Serial Search** to display devices with serial numbers starting with number sequence entered.

4. Click **Select**.
5. If required, enter a start and end date for the report.
The start date defaults to the earliest entry in the database and the end date defaults to today's date.
6. Click **Create Report**.
The report is generated and displayed.

Export as an Excel or PDF file. Delete the report.

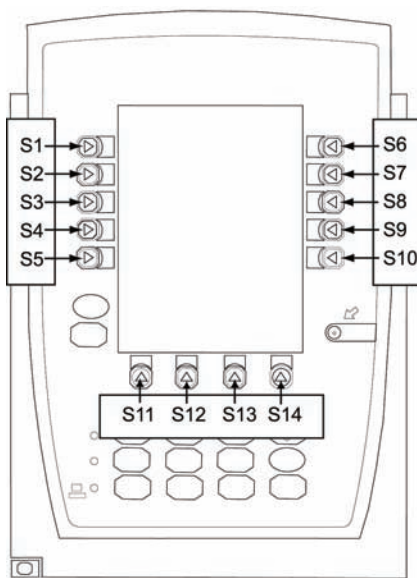
Generated reports are added to the **Report** list. Click the arrows in the column headings to sort the report by column. Shift-click a second heading to choose it as a secondary sort column.

Logs

This section describes how to read the System Maintenance Software logs. There are three types of logs:

- **Battery Log:** Contains internal and external events relating to the battery, such as the charging state and other battery status conditions.
- **Event Log:** Contains a complete record of operating events, such as keyboard presses, alarms, module attachments, external communications, and infusion starts and stops. The log might also contain additional information in coded form about changes that occurred on the display screen and other internal software events.
- **Error Log:** Contains a record of any abnormal events, such as internal software and hardware malfunctions.

System Maintenance Software logs refer to the soft keys on the PC Unit:



Log Definitions

System Maintenance Software logs can include these entries:

Log Definitions

Log Entry	Definition
AIL	Air-in-Line
AIU	PC Unit
AKB	(AIU) PC Unit keyboard (keypad) processor
ATOD	Analog to digital converter
ATTACHED_UNIT_FAULT	A malfunction from an attached module, also logged in PC Unit error log
AVA	Audio/visual event

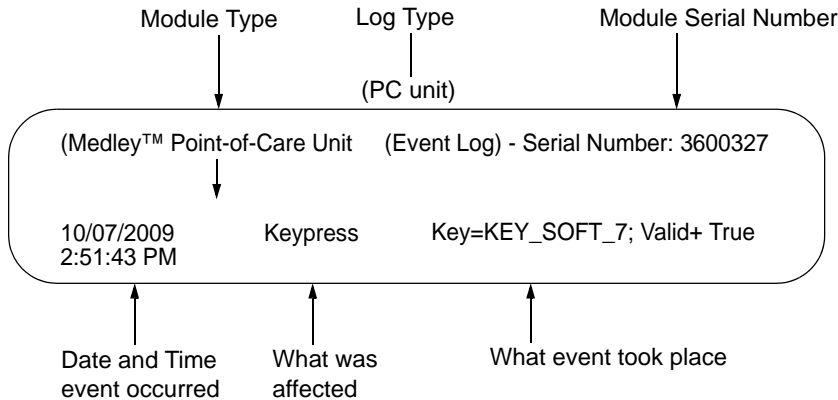
Log Definitions (Continued)

Log Entry	Definition
BFD	Big file download
BLOCK_STEP_COMPLETE	Indicates an infusion step (delay, bolus, keep vein open [KVO], and so on) is complete
CALLSTACK_DATA	Software execution history logged for debugging purposes
CBIT	Continuous built-in test
CIB, CI	Communication Board
COMM	Communications
CPU	Central processing unit
DATASET_I_D	The ID of the loaded Data Set
DEBUG_DATA / DEBUG_EVENT	Miscellaneous data logged for debugging purposes
DRAM	Dynamic RAM
EMI	Electromagnetic interference
ETC	EtCO ₂ Module
EXT_RAM	External RAM
FD	Fluid delivery domain
FORM_REQUEST	A new page or combination of pages is displayed on the PC Unit
INF	Infusion
INT_RAM	Internal RAM
INVALID_TRANSITION_FAULT	Illegal state machine event
KEY_0, KEY_1, etc	Numeric or other "hard-labeled" keys on device
KEY_SOFT_1	Software-labeled or "soft" keys whose meaning is determined by which page is displayed
LB3	Low Battery Event 3: all devices pause, low battery alert
LB4	Low Battery Event 4: battery completely discharged, system powers down, backup alarm
LTI	Log table invariant, a tag used to mark special data, such as end of a log
LVP	Pump Module
MALFUNCTION	Hardware error
MMS	Medication Management System
NET	Network
OEM	Original equipment manufacturer
OS	Operating system

Log Definitions (Continued)

Log Entry	Definition
PATIENT_ASSOC	A patient associated with PC Unit
PATIENT_I_D	Patient identification
PM	Preventive maintenance
PCU	PC Unit. The central programming and power supply module of the Alaris® System
POST	Power on self test
PROGRAM	A complete sequence of infusion steps, including delay, primary, secondary, bolus, KVO, and so on
PROGRAM_COMPLETE	Indicates all programmed steps are complete
PSP	Power supply processor
PUMP	Infusion component of Alaris® System
PWR	Power domain
RAM	Random access memory
ROM	Read-only memory
RTC	Real-time clock
SAFETY_SWITCH	An element that can turn off power to motor any time
SHUTDOWN_COMPLETE	Shutdown tasks complete, ready to power down
SKB	SpO ₂ Module keyboard processor
SOFTWARE_FAULT	Software error
SPO, SP_O2	SpO ₂ Module (pulse oximetry module of Alaris® System)
SPR	Safety processor
SRAM	Static RAM
SS	Subsystem
STEP	A single infusion step (delay, primary, secondary, bolus, KVO, and so on)
STRING	General event for a log string
SYRINGE_SELECTION	User selected syringe
UI	User interface
UNINIT_INTERRUPT_FAILURE	Un-initialized interrupt failure, an illegal interrupt occurrence
VPMR	Volume per mechanical revolution
VTBI	Volume to be infused
WD	Watchdog

Example Log



The following table shows an example Event log. In this example:

- two Pump Modules are attached to the PC Unit (Pump Module A and Pump Module B).
- Pump Module B is set to run at a rate of 125 ml/h and VTBI of 125 ml.
- the Pump Module door opened, causing an alarm, and then it closed.
- the Pump Module is restarted and then turned off.

An explanation of each log entry is in shaded text.

Example Log

Log Time	Category ID	Details
10/11/2009 7:27:16 AM	POWERED_ON	ModelNumber=8000; SerialNumber=3600327; SwVersionNumber=1.10.2.0
Device powered on. Serial number and software version of PC Unit.		
10/11/2009 7:27:17 AM	PUMP_IDLE	Serial Number=3647764
Pump Module is not running. Pump Module serial number.		
10/11/2009 7:27:17 AM	UNIT_ADDED	Model Number=8100; Serial Number=3647764; UnitLabel=A
Module has responded to PC Unit. First Pump Module detected is designated as module A.		
10/11/2009 7:27:17 AM	PUMP_IDLE	SerialNumber=3648579
Pump Module is not running. Pump Module serial number.		
10/11/2009 7:27:17 AM	UNIT_ADDED	ModelNumber=8100; SerialNumber=3648579; UnitLabel=B
Module has responded to PC Unit. Second Pump Module detected is designated as module B.		
10/11/2009 7:27:17 AM	FORM_REQUEST	Form=NEW_PATIENT; FormRequest=FORM_REQUEST
NEW PATIENT? screen is displayed on PC Unit.		

Example Log (Continued)

Log Time	Category ID	Details
10/11/2009 7:27:18 AM	AVA_EVENT	SourceType=POWER_MANAGEMENT; SourceContext=0; EventType=PWR_BATTERY_OK
Audio/visual event: PC Unit battery power check OK.		
10/11/2009 7:27:19 AM	PUMP_IDLE	SerialNumber=3647764
Pump Module is not running. Serial number identifies module A.		
10/11/2009 7:27:19 AM	PUMP_IDLE	SerialNumber=3648579
Pump Module is not running. Serial number identifies module B.		
10/11/2009 7:27:21 AM	KEYPRESS	Key=KEY_SOFT_7; Valid=True
Key press on PC Unit: soft key 7. (Soft key 7 on NEW PATIENT? screen corresponds to No.)		
10/11/2009 7:27:24 AM	UNIT_KEYPRESS	DeviceAddress=3648579; Key=KEY_UNIT_SELECT; Valid=True
Key pressed on module: serial number identifies Pump Module B CHANNEL SELECT key.		
10/11/2009 7:27:24 AM	GUARDRAIL_START	SerialNumber=3648579; ProfileIndex=255; IsAnesthesiaMode=False
Guardrails® software starts, performs check on Pump Module B (identified by serial number).		
10/11/2009 7:27:24 AM	FORM_REQUEST	Form=BASIC_INFUSION; FormRequest=FORM_REQUEST
PC Unit displays Infusion Setup screen for most recently identified Pump Module (module B).		
10/11/2009 7:27:26 AM	KEYPRESS	Key=KEY_SOFT_1; Valid=True
Key press on PC Unit: soft key 1. (Soft key 1 on Infusion Setup screen corresponds to RATE.)		
10/11/2009 7:27:27 AM	KEYPRESS	Key=KEY_1; Valid=True
Key press on PC Unit: numeric key 1 (RATE is 1).		
10/11/2009 7:27:28 AM	KEYPRESS	Key=NUM_2; Valid=True
Key press on PC Unit: numeric key 2 (RATE is 12).		
10/11/2009 7:27:28 AM	KEYPRESS	Key=KEY_5; Valid=True
Key press on PC Unit: numeric key 5 (RATE is 125).		
10/11/2009 7:27:29 AM	KEYPRESS	Key=KEY_SOFT_2; Valid=True
Key press on PC Unit: soft key 2 (soft key 2 on Infusion Setup screen corresponds to VTBI).		

Example Log (Continued)

Log Time	Category ID	Details
10/11/2009 7:27:29 AM	KEYPRESS	Key=KEY_1; Valid=True
Key press on PC Unit: numeric key 1 (VTBI is 1).		
10/11/2009 7:27:30 AM	KEYPRESS	Key=NUM_2; Valid=True
Key press on PC Unit: numeric key 2 (VTBI is 12).		
10/11/2009 7:27:30 AM	KEYPRESS	Key=KEY_5; Valid=True
Key press on PC Unit: numeric key 5 (VTBI is 125).		
10/11/2009 7:27:33 AM	KEYPRESS	Key=KEY_SOFT_14; Valid=True
Key press on PC Unit: soft key 14 (soft key 14 on Infusion Setup screen corresponds to START).		
10/11/2009 7:27:33 AM	PROGRAM_DATA	ProgramID=1; ProgramType=PRIMARY_INFUSION; NumSteps=2
PC Unit program: primary infusion.		
10/11/2009 7:27:33 AM	BLOCK_STEP_DATA	StepNum=0; Rate=125; VTBI=125; InfuseAll=False; FracRate=False
Infusion step: rate=125, VTBI=125.		
10/11/2009 7:27:33 AM	BLOCK_STEP_DATA	StepNum=1; Rate=1; VTBI=0; InfuseAll=True; FracRate=False
Infusion step: when primary infusion is complete, KVO to run at rate=1, VTBI=0.		
10/11/2009 7:27:33 AM	GUARDRAIL_END	Type=0
Guardrails® software check complete: no problems detected.		
10/11/2009 7:27:33 AM	PROGRAM_START	SerialNumber=3648579; ProgramID=1; ProgramType=PRIMARY_INFUSION
Program start: primary infusion on Pump Module B.		
10/11/2009 7:27:33 AM	BLOCK_STEP_DATA	StepNum=0; Rate=125; VTBI=125; InfuseAll=False; FracRate=False
Infusion step: rate=125, VTBI=125.		
10/11/2009 7:27:33 AM	FORM_REQUEST	Form=BASIC_INFUSION; FormRequest=CANCEL_FORM
PC Unit displays screen: basic infusion screen displays modules operating and VTBI of each (in this example, only Pump Module B is operating, VTBI =125).		
10/11/2009 7:27:34 AM	PUMP_INFUSING	SerialNumber=3648579
Pump infusing: Pump Module B.		

Example Log (Continued)

Log Time	Category ID	Details
10/11/2009 7:27:35 AM	AVA_EVENT	SourceType=INFUSOR; SourceContext=3648579;EventType=INFUSOR_START ED
Audio/visual event: Pump Module B starts (INFUSING indicator lights).		
10/11/2009 7:27:40 AM	PUMP_ALARMED	SerialNumber=3648579
Pump Module alarm: Pump Module B (ALARM indicator lights, pump stops running).		
10/11/2009 7:27:40 AM	AVA_EVENT	SourceType=INFUSOR; SourceContext=3648579;EventType=INFUSOR_DOOR_ OPENED
Audio/visual event: Pump Module B door open.		
10/11/2009 7:27:43 AM	PUMP_ALARMED	SerialNumber=3648579
Pump Module alarm: Pump Module B.		
10/11/2009 7:27:43 AM	AVA_EVENT	SourceType=INFUSOR; SourceContext=3648579; EventType=INFUSOR_DOOR_CLOSED
Audio/visual event: Pump Module B door closed.		
10/11/2009 7:27:44 AM	UNIT_KEYPRESS	DeviceAddress=3648579; Key=KEY_PUMP MODULE_RESTART; Valid=True
Module key press: Pump Module B RESTART key.		
10/11/2009 7:27:44 AM	PUMP_INFUSING	SerialNumber=3648579
Pump Module infusing: Pump Module B.		
10/11/2009 7:27:45 AM	AVA_EVENT	SourceType=INFUSOR; SourceContext=3648579; EventType=INFUSOR_STARTED
Audio/visual event: Pump Module B starts (INFUSING indicator lights).		
10/11/2009 7:28:04 AM	UNIT_KEYPRESS	DeviceAddress=3648579; Key=KEY_PUMP MODULE_PAUSE; Valid=True
Module key press: Pump Module B PAUSE key.		
10/11/2009 7:28:04 AM	PUMP_PAUSED	SerialNumber=3648579
Pump Module paused: Pump Module B.		
10/11/2009 7:28:04 AM	AVA_EVENT	SourceType=INFUSOR; SourceContext=3648579; EventType=INFUSOR_PAUSED
Audio/visual event: Pump Module B STANDBY indicator lights.		
10/11/2009 7:28:41 AM	UNIT_KEYPRESS	DeviceAddress=3648579; Key=KEY_UNIT_CHANNEL_OFF; Valid=True
Module key press: Pump Module B CHANNEL OFF key.		

Example Log (Continued)

Log Time	Category ID	Details
10/11/2009 7:28:42 AM	PUMP_IDLE	SerialNumber=3648579
Pump Module is not running: Pump Module B.		
10/11/2009 7:28:42 AM	PROGRAM_COMPLETE	SerialNumber=3648579; ProgramID=1; ProgramType=PRIMARY_INFUSION
Program complete: primary infusion has stopped on Pump Module B.		
10/11/2009 7:28:43 AM	AVA_EVENT	SourceType=INFUSOR; SourceContext=3648579; EventType=INFUSOR_STOPPED
Audio/visual event: Pump Module B stopped.		
10/11/2009 7:28:43 AM	FORM_REQUEST	Form=POWERING_DOWN; FormRequest=FORM_REQUEST
PC Unit display: Flashes a "Powering Down" message for about three seconds.		

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Glossary

A | C | D | E | F | I | N | P | S | T | W

A

Alaris® Server

The server that manages wireless communication with Alaris® products.

ASM

Alaris® System Maintenance

Auto-ID Module

The bar code component (Model 8600) of the Alaris® System.

C

communication (comm) board (CB)

An optional wireless network card (sometimes integrated in the motherboard). It is installed in the PC Unit and allows communication with the Alaris® Server.

D

Data Set

Created using the Editor Software authoring tool and then transferred to the PC Unit. A Data Set reflects the facility's best-practice guidelines for IV drug administration and includes: Profile Drug Libraries, Clinical Advisories, instrument configurations, and channel Label Libraries.

delimiter

A character that marks the beginning or end of a unit of data.

DFU

Directions for Use.

E

EtCO₂ Module

The capnograph component (Model 8300) of the Alaris® System.

F

flash

The process of electronically reimaging the Operating System.

I

IUI Connector

The inter-unit interface connector that connects the PC Unit and modules.

N

network configuration package

A package that contains up to eight WSec (wireless security) Network Profiles.

P

PCA Module

The Patient Controlled Analgesia (PCA) component (Model 8120) of the Alaris® System.

PC Unit

The central programming, monitoring, and power supply component of the Alaris® System.

preventive maintenance

The set of tasks that are performed a minimum of once a year to ensure proper functioning of a module.

Pump Module

A large volume infusion component (Model 8100) of the Alaris® System.

S

SpO₂ Module

The pulse oximetry components (Models 8210 and 8220) of the Alaris® System.

Syringe Module

The syringe infusion component (Model 8110) of the Alaris® System.

T

transfer

Copy hospital-created data from the personal computer to a PC Unit.

W

wireless card

The connection device that allows the Alaris® Server to communicate with the communications board on a PC Unit.

WSec

Wireless security.

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