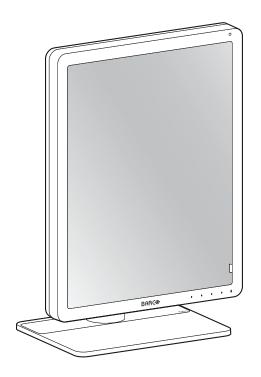
Nio 2MP LED Display



User Guide

MDNC-2221



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Welcome!

1

1.1 What's in the box

Overview

- MDNC-2221 display
- · User guide
- · System sheet
- Documentation disc
- DisplayPort cable
- DVI cable
- · USB cable
- Mains cable(s)
- · External power supply

If you ordered a Barco display controller, it's also in the box together with its accessories. A dedicated user guide is available on the documentation disc.



Keep your original packaging. It is designed for this display and is the ideal protection during transport and storage.

1.2 At a glance

Front

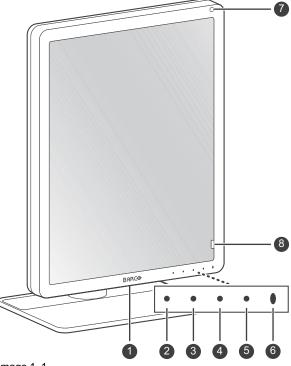
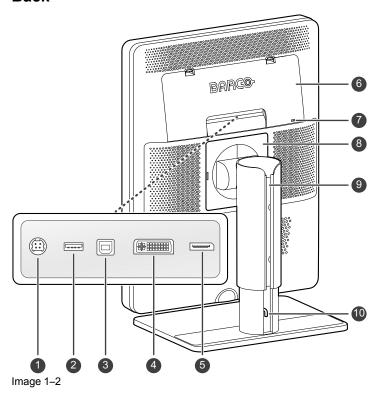


Image 1-1

- 1. USB-A 2.0 downstream connector
- 2. Left key
- 3. Right key
- 4. Menu key
- 5. Standby key
- 6. Power status LED
 - Off: Display not powered, or display operational but power LED function disabled in OSD (see "Power status LED", page 22)
 - Steady green: Display operational

- Steady amber: Display switched off via the standby key, or display in hibernate mode (requires DPMS mode and Hibernate to be enabled in the OSD menu, see "DPMS mode", page 23 and "Hibernate", page 23)
- 7. Ambient light sensor
- 8. Front sensor

Back



- 1. +24 VDC power input
- 2. USB-A 2.0 downstream connector
- 3. USB-B 2.0 upstream connector
- 4. DVI video input
- 5. DisplayPort video input
- 6. Connector compartment cover
- 7. Kensington security lock
- 8. VESA mount cover plate
- 9. Cable duct
- 10. Foot lock clip

Welcome!

Installation

2

2.1 Display controller installation

About

Before you install your display and connect it with the workstation, make sure to have a suitable¹ display controller installed in the workstation.



The Nio 2MP LED Display display operates at its full specifications when driven by a Barco **MXRT display controller** and **MXRT driver** (which also includes the Barco **Intuitive Workflow Tools**). If you ordered a MXRT display controller, it is included in the box of your display.

For more information and installation instructions, please check the Barco "Display Controller and Intuitive Workflow Tools" user guide on the included documentation CD, or at www.barco.com/support.

2.2 Display position adjustment

To adjust the display position

1. Pull out the red foot lock clip from the hole at the back of the foot.

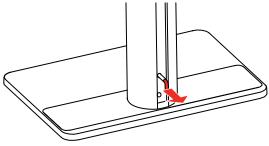


Image 2-1

2. You can now safely tilt, swivel, pivot, raise and lower the display as desired.

For a list of compatible display controllers, please refer to the compatibility matrix available on my.barco.com (MyBarco > My Support > Healthcare > Compatibility Matrices > Barco Systems Compatibility Matrices)

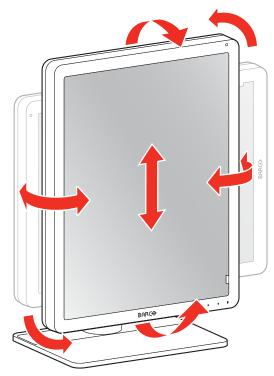
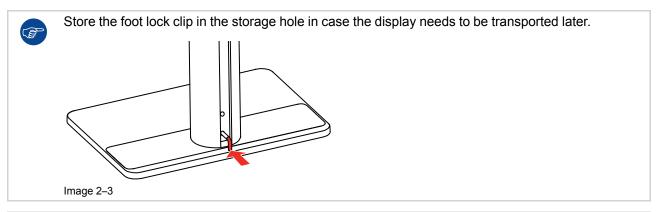


Image 2-2



WARNING: The display must be in its highest position before it can be properly pivoted.



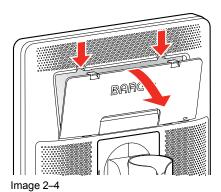


If, after installing the display of the system, you change the panel orientation while an image is on the screen, the result depends on the graphic board and the resolution of the image. In some cases the image will be rotated automatically, in other cases it will not be rotated (e.g., when pixels would be lost after rotation). If necessary, change the image resolution in the display control panel and restart the system after changing the orientation.

2.3 Cable connections

To connect the cables

1. Open the connector compartment. Do this by gently pushing the two lips on top of the cover, after which it can be removed from the display.



2. Connect a video source from your workstation to the corresponding video inputs on the display. **Caution:** Connect only one of the two video inputs. Connecting both inputs simultaneously will result in driver errors.

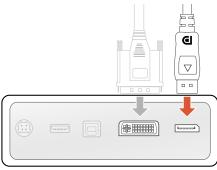


Image 2-5

3. Connect the workstation's USB host to the USB upstream connector on the display if you want to make use of QAWeb or any of the display USB downstream connectors (e.g. to connect a keyboard, mouse, touch pad or other peripheral).

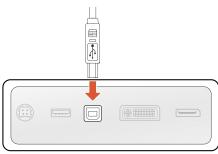


Image 2-6

4. Connect the supplied external DC power supply to the power input on the display.



Image 2-7

5. Route all cables through the cable duct in the stand of your display. For better strain-relief and shielding of the cables, fix the cables with the cable straps at the inside of the connector compartment.

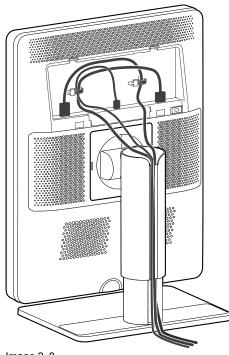


Image 2-8

6. Close the connector compartment cover. You will hear a "click" sound when the cover is in position.

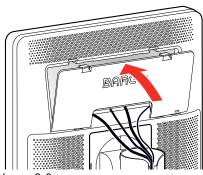


Image 2-9

7. Connect the external DC power supply to a **grounded** power outlet by using one of the power cables included with the display.

2.4 MXRT driver and Intuitive Workflow Tools installation

About

When you are using a Barco MXRT display controller, you can start up your Nio 2MP LED Display display system and install the **MXRT driver** and **Intuitive Workflow Tools**. Barco's Intuitive Workflow Tools are designed to increase visibility of subtle details, improve focus during reading sessions, and accelerate workflow.

To install the MXRT driver and Intuitive Workflow Tools

- 1. Switch on your Nio 2MP LED Display as described in "Standby switching", page 19.
- 2. Turn on the workstation connected to your display.
 Your display will be running in a basic video mode at a default refresh rate when first time starting up.
- 3. Download the latest MXRT driver and Intuitive Workflow Tools from www.barco.com/mxrt.

- Install the MXRT driver and Barco's Intuitive Workflow Tools as described in the Barco "Display Controller and Intuitive Workflow Tools" user guide on the included documentation CD, or at www.barco.com/support.
- 5. When the drivers are completely installed, your display will automatically detect the connected video input signal(s) and apply the correct video mode and refresh rate.

2.5 QAWeb registration

About

QAWeb Enterprise helps you manage quality and assure compliance of your expanding healthcare enterprise with less effort, lower cost, and complete confidence. This fully automated and secure system guarantees a consistent image quality and uptime for all registered imaging display systems within your facility and across your enterprise. Learn more at www.barco.com/qaweb.

To register your display system to your QAWeb Enterprise organization, the QAWeb Enterprise Agent must be installed and running on your workstation and it must be able to communicate with the QAWeb Enterprise cloud service.

For more information and instructions, check the QAWeb Enterprise user guide or www.barco.com/support/gaweb-enterprise.



When still using the older MediCal QAWeb, visit <u>www.barco.com/support/medical-qaweb</u> to obtain the installation package and user documentation.

2.6 VESA-mount installation



CAUTION: Use suitable mounting apparatus to avoid risk of injury.



WARNING: Never move a display attached to an arm by pulling or pushing the display itself. Instead, make sure that the arm is equipped with a VESA compliant handle and use this to move the display.

Please refer to the instruction manual of the arm for more information and instructions.



WARNING: Use a mount that is compliant with the VESA 100 mm standard.

Use a mount that can support the weight of the display. Refer to the technical specifications of this display for the applicable weight.

Overview

The panel, standard attached to the tilt & swivel foot, is compatible with the VESA 100 mm standard. So it can be used with an arm stand according to the VESA 100 mm standard.

Therefore, the tilt & swivel foot must be removed from the panel.

1. Put the display in the lowest position and lock the height mechanism.

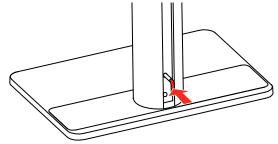
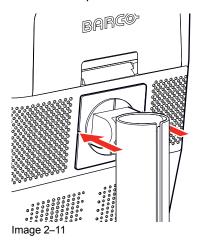


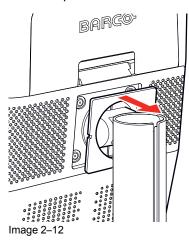
Image 2-10

2. Put the display face down on a clean and soft surface. Be careful not to damage the panel screen.

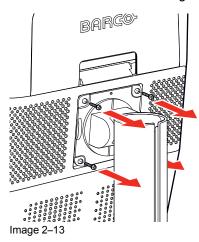
3. Remove the plastic cover with a flathead screw driver.



4. Slide the plastic cover over the neck of the foot.



5. Remove the four screws fixing the foot while supporting the foot.



6. Attach the arm stand **firmly** to the panel using 4 screws M4 x 8 mm.

Installation

Daily operation

3

3.1 Recommendations for daily operation

Optimize the lifetime of your display

Enabling the Display Power Management System (DPMS) of your display will optimize its diagnostic lifetime by automatically switching off the backlight when the display is not used for a specified period of time. By default, DPMS is enabled on your display, but it also needs to be activated on your workstation. To do this, go to the "Power Options" of your workstation.



Barco recommends setting DPMS activation after 20 minutes of non-usage.

Use a screen saver to avoid image retention

Prolonged operation of an LCD with the same content on the same screen area may result in a form of image retention.

You can avoid or significantly reduce the occurrence of this phenomenon by using a screen saver. You can activate a screen saver in the "Display properties" window of your workstation.



Barco recommends setting screen saver activation after 5 minutes of non-usage. A good screen saver displays moving content.

In case you are working with the same image or an application with static image elements for several hours continuously (so that the screen saver is not activated), change the image content regularly to avoid image retention of the static elements.

Understand pixel technology

LCD displays use technology based on pixels. As a normal tolerance in the manufacturing of the LCD, a limited number of these pixels may remain either dark or permanently lit, without affecting the diagnostic performance of the product. To ensure optimal product quality, Barco applies strict selection criteria for its LCD panels.

Enhance user comfort

Every Barco multi-head display system is color matched to the tightest specifications in the market.



Barco recommends keeping color-matched displays together. Furthermore, it is important to use all displays of a multi-head configuration at the same rate to preserve color matching throughout the economic lifetime of the system.

Maximize quality assurance

QAWeb Enterprise helps you manage quality and assure compliance of your expanding healthcare enterprise with less effort, lower cost, and complete confidence. This fully automated and secure system guarantees a consistent image quality and uptime for all registered PACS display systems within your facility and across your enterprise.



Barco highly recommends to use QAWeb Enterprise. Learn more at www.barco.com/qaweb.

3.2 Key indicator lights

About the key indicator lights

By default, the indicator lights of the keys will be off which makes the keys unavailable at that moment. To make the keys illuminate and available for further actions touch one of the keys. As a result, all keys will be illuminated and are now available for further actions. However, if no further actions are taken within the following 5 seconds, the keys will dim again.



The key auto-dim function can be disabled in the OSD menus. Please refer to "Key indicator lights", page 18 for detailed instructions on how to do this.

3.3 Standby switching

About standby switching



The connected power supply also provides a switch that can be used to turn the power completely off. To use the display, please make sure to switch on this power supply. This can be done by pushing the on/off switch on the power supply into the "|" position.

Switching on the display while it is in standby mode or vice versa can be done by:

- 1. Illuminate the keys as previously described.
- 2. While the keys are illuminated, touch the standby key for approximately 2 seconds.

As a result, the display will switch on or will switch to standby mode.



In case of a power outage recovery, your display will always start-up in the power mode it was in before the power interruption (i.e. standby or on). This protects your display against inadvertent image retention problems.

3.4 OSD menu access

About the OSD menu

The OSD menu allows you to configure different settings to make your Nio 2MP LED Display fit your needs within your working environment. Also, you can retrieve general information about your display and its current configuration settings through the OSD menu.

To access the OSD menu

- 1. Press any of the front keys to illuminate and activate them.
- 2. While the keys are illuminated, press the Menu key (). The OSD main menu comes up.
- 3. If the OSD menu locked message appears, wait for the message to disappear, then press the following key sequence: Menu Left Right Left Menu. After this, the OSD main menu comes up.

3.5 OSD menu navigation

To navigate through the OSD menus

Navigating through the OSD menus can be done by:

- Use the left/right keys to move through the (sub)menus, change values or make selections.
- To go into a submenu or confirm adjustments and selections, use the menu key.
- Use the standby key to cancel adjustments or exit a (sub)menu.
- Exit all OSD menus at once by touching the standby key for approximately 2 seconds.

The key icons are displayed above the keys and are adapted to the function that it is used for (menu dependent):



Left, Right

Menu

Daily operation



Enter



Cancel



Standby (IEC 60417-5009)

Advanced operation

4

4.1 OSD menu language

About the OSD menu language

By default, the OSD menu comes up in English. However, there's a wide range of other languages available for the OSD menu of your Nio 2MP LED Display.

To change the language of the OSD menu:

- 1. Bring up the OSD main menu.
- 2. Navigate to the Configuration > User Interface > Menu menu.
- 3. Enter the Language submenu.
- 4. Select one of the available languages and confirm.

4.2 OSD menu automatic close function

About the OSD menu automatic close function

By default, the OSD menu will disappear automatically after approximately 90 seconds of inactivity. However, this function can be disabled so that the OSD menu remains on the screen until manually closed.

To enable/disable the OSD menu automatic close function:

- 1. Bring up the OSD main menu.
- 2. Navigate to the Configuration > User Interface > Menu menu.
- 3. Enter the Automatic Close submenu.
- 4. Select Enabled/Disabled as desired and confirm.

4.3 Power status LED

About the power status LED

By default, when the display is switched on, the power status LED is dimmed. This behavior can be changed so that the power status LED will be **green** when the display is switched on.



When the display is in stand-by mode, the power status LED will always turn orange, even when the power status LED is disabled.

To enable/disable the power status LED:

- 1. Bring up the OSD main menu.
- 2. Navigate to the Configuration > User Interface > Indicator Lights menu.
- 3. Enter the Power Status submenu.
- 4. Select Enabled/Disabled as desired and confirm.

4.4 Key indicator lights

About the key indicator lights

By default, after lighting up, the key indicator lights will dim again if no further actions are taken within the following 5 seconds. However, this behavior can be changed so that the key indicator lights are always on or always off.

To configure the key indicator lights

- 1. Bring up the OSD main menu.
- 2. Navigate to the Configuration > User Interface > Indicator Lights menu.
- 3. Enter the Keys submenu.

4. Select Automatic/Always On/Always Off as desired and confirm.

4.5 Power lock function

About the power lock function

By enabling the power lock function, the Nio 2MP LED Display is forced to remain switched on. This means that it can't be switched to stand-by mode manually until the power lock function is disabled again.

To enable/disable the power lock function:

- 1. Bring up the OSD main menu.
- 2. Navigate to the Configuration > User Interface > Controls menu.
- 3. Enter the Power Lock submenu.
- 4. Select Enabled/Disabled as desired and confirm.

4.6 DPMS mode

About DPMS mode

Enabling the Display Power Management System (DPMS) mode on your display will optimize its diagnostic lifetime by automatically switching off the backlight when the display is not used for a specified period of time. By default, DPMS mode is enabled on your display, but it also needs to be activated on your workstation. To do this, go to the "Power options properties" window of your workstation.



Barco recommends setting DPMS activation after 20 minutes of non-usage.



When DPMS mode is enabled, an additional power saving function becomes available: hibernate. See "Hibernate", page 23 for more information.

To enable/disable DPMS mode on your display:

- 1. Bring up the OSD main menu.
- 2. Navigate to the Configuration > Power Management menu.
- 3. Enter the DPMS Mode submenu.
- 4. Select Enabled/Disabled as desired and confirm.

4.7 Hibernate

About hibernate

When hibernate is enabled, not only the backlight will be switched off, but also other functionalities will be disabled to further reduce power consumption to a minimum. This happens after a specific period of time which can be manually adjusted.



Hibernate can only be enabled on your display when the DPMS mode is enabled first. Therefore, please refer to "DPMS mode", page 23 to do this.



Please connect your keyboard or mouse to your PC rather than to the display's USB ports when hibernate is enabled.

To enable/disable hibernation on your display:

- 1. Bring up the OSD main menu.
- 2. Navigate to the Configuration > Power Management menu.
- 3. Enter the Hibernate submenu.

4. Select Enabled/Disabled as desired and confirm.

To specify the hibernate time-out:

- 1. Bring up the OSD main menu.
- 2. Navigate to the Configuration > Power Management menu.
- 3. Enter the Hibernate Timeout submenu.
- 4. Set the time-out value as desired and confirm.

4.8 Luminance target

About the luminance target

The luminance target of your Nio 2MP LED Display is adjustable over a predefined range. When you change the luminance target, the display will adjust its backlight to reach the target.

To set the luminance target:

- 1. Bring up the OSD main menu.
- 2. Navigate to the *Configuration > Calibration* menu.
- 3. Enter the Luminance Target submenu.
- 4. Set a luminance target value as desired and confirm.



The default, factory DICOM calibrated luminance value is available in the technical specifications table. The guaranteed backlight lifetime is valid for this setting.

4.9 Viewing modes

About viewing modes

The Nio 2MP LED Display can be used in two viewing modes:

- **Diagnostic:** This mode provides the full calibrated luminance and is intended for using the display for diagnostic purposes.
- **Text:** In this mode, the luminance is reduced to approximately half of the luminance. This is intended for using the display with office applications such as word processing.
 - Please note that text mode is not persistent, once powered off, the unit will restart in diagnostic mode.



The diagnostic mode should always be selected when the Nio 2MP LED Display is intended to be used in a diagnostic environment.

To select a viewing mode:

- 1. Bring up the OSD main menu.
- 2. Navigate to the Configuration > Calibration menu.
- 3. Enter the Viewing Mode submenu.
- 4. Select Diagnostic/Text as desired and confirm.

4.10 Display functions

About display functions

Native, uncorrected panels will display all grayscale/color levels with luminance increments that are not optimal for crucial diagnostic information. Studies have shown however, that in medical images certain grayscale/color parts contain more diagnostic information then others. To respond to these conclusions,

display functions have been defined. These functions emphasize on these parts containing crucial diagnostic information by correcting the native panel behavior.

The available display functions for your Nio 2MP LED Display are:

- Native: If you select Native, the native panel behavior will not be corrected.
- **Dynamic Gamma 1.8** or **2.2**: These are gamma functions that are shifted to take into account the non-zero luminance of an LCD panel when driven with a "black" signal. They are especially useful in CT applications to improve the perception of low Hounsfield values.
- DICOM: DICOM (Digital Imaging and Communications in Medicine) is an international standard that was
 developed to improve the quality and communication of digital images in radiology. In short, the DICOM
 display function results in more visible grayscales in the images. Barco recommends selecting the DICOM
 display function for most medical viewing applications.
- User: This display function will be automatically selected when display functions are defined by MediCal QAWeb.
- **Gamma 1.8** or **2.2**: Select one of these display functions in case the display is to replace a CRT display with a gamma of 1.8 or 2.2 respectively.



The settings of the display must be adapted to suit the requirements of the visualization software. In case of doubt, please contact the vendor of the visualization software.

To select a display function:

- 1. Bring up the OSD main menu.
- 2. Navigate to the Configuration > Calibration menu.
- 3. Enter the Display Function submenu.
- 4. Select one of the available display functions and confirm.

4.11 Reading rooms

About reading rooms



Reading rooms can only be selected when the DICOM display function is selected. Therefore, please refer to "Display functions", page 24 to correctly set the display function.

The American Association of Physicists in Medicine (AAPM) composed a list of pre-defined reading rooms. Each of these reading rooms are defined by following parameters:

· the maximum light allowed in this type of room

The available reading rooms for your Nio 2MP LED Display are:

- **Dark Room:** Corresponds to light conditions of ideal dark room. This setting disables the ambient light compensation.
- **CR/DR/ MAMMO:** Corresponds to light conditions in diagnostic reading rooms for computed radiology, digital radiology or mammography. This setting has the lowest maximum ambient light.
- **CT/MR/NM:** Corresponds to light conditions in diagnostic reading rooms for computed tomography, magnetic resonance or nuclear medicine scans.
- Staff Office: Corresponds to light conditions in office rooms.
- Clinical Viewing Room: Corresponds to light conditions in diagnostic reading rooms for clinical viewing.
- **Emergency Room:** Corresponds to light conditions in emergency rooms.
- **Operating Room:** Corresponds to light conditions in operating rooms. This setting has the highest maximum ambient light.

To select a reading room:

- 1. Bring up the OSD main menu.
- 2. Navigate to the Configuration > Calibration > Reading Room menu.
- 3. Enter the *Reading Room* submenu.
- 4. Select one of the available reading rooms and confirm.

4.12 Display orientation

About Display orientation

Your display automatically detects its physical orientation (portrait or landscape) and, by default, automatically adjusts the image orientation to this. This means that when your display is physically rotated, the image will rotate along.

The OSD menu however, allows to overrule this behavior and force the image orientation to portrait or landscape regardless of the physical orientation of the display. This may be especially useful when operating your display with the screen facing upwards.

To set the Display orientation

- 1. Bring up the OSD main menu.
- 2. Navigate to the Configuration > Image Sources menu.
- 3. Enter the Display Orientation submenu.
- 4. Select Landscape/Portrait/Automatic as desired and confirm.

4.13 Video input signals

About input signals

The available input signals for your display are:

- DisplayPort 1: The input corresponding to the DisplayPort connector.
- DVI 1: The input corresponding to the DVI connector.
- Automatic Selection: The input is automatically selected.

To manually select a video input signal:

- 1. Bring up the OSD main menu.
- 2. Navigate to the Configuration > Image Source menu.
- 3. Enter the Input Signal submenu.
- 4. Select one of the available input signals and confirm.

4.14 EDID timings

About EDID timings

Following EDID timings are available for your Nio 2MP LED Display:

- Refresh Rate: Allows to manually select the refresh rate of the image source video input signal depending on the maximum refresh rate of the display controller connected to your display.
- Preferred Orientation: Allows to change the orientation of the image source video input signal to landscape, portrait or to let the display automatically assign the correct orientation.
- Color Depth: Allows to change the color depth to 8 or to 10 bit.

To manually set EDID timings:

- 1. Bring up the OSD main menu.
- 2. Navigate to the Configuration > Image Source menu.
- 3. Enter the *Timings* submenu.
- 4. Select Refresh Rate, Preferred Orientation or Color Depth.
- 5. Select one of the available settings and confirm.

4.15 Display info

About display info

Your display serial number, color type, native resolution, firmware versions, etc. are available in a dedicated submenu of the OSD menu.

To retrieve info about your display:

- 1. Bring up the OSD main menu.
- 2. Navigate to the About this Display menu to make the information visible on the screen.

4.16 Display status

About display status

The Status submenu of the OSD menu provides info on the current status of your display (runtimes, temperatures, etc.), the status of the connected image sources (video encoding mode, timings, etc.), the current calibration status of your display (display function, luminance, ALC, etc.) and the status about activated connections.

To retrieve the status of your display:

- 1. Bring up the OSD main menu.
- 2. Navigate to the Status menu.
- 3. Enter the Display, Image Sources, Calibration or Connectivity submenu as desired.

Advanced operation

Maintenance

5.1 Scheduled maintenance

About

The Nio 2MP LED Display does not require any scheduled maintenance or calibration activities. We recommend to use QAWeb with the Barco default tests and frequencies to calibrate and maintain the display, or to return the display to a Barco approved maintenance organization. In any case of doubts, please contact Barco Healthcare.

5.2 Cleaning instructions

To clean the display

Apply a cleaning/disinfecting product to a soft lint-free cloth, such as a microfiber or gauze and rub the display surface thoroughly. In order to be effective, all surfaces must be cleaned for a certain amount of time (ranging from 30 seconds to 2 minutes).

Use a cleaning/disinfecting product that is alcohol-, alkali-, water- or chlorine-based. Common examples are:

- Isopropanol 100%
- Ethanol 70%
- 0.5% Chlorehexidine in 70% ethanol/isopropanol
- Ortho-Phthalaldehyde (OPA) 0.55%
- · Haemo-sol, 1% in water
- · 250 ppm Chlorine solution
- 1.0% lodine in 70% ethanol
- 1.6% aqueous ammonia
- "Green soap" (USP)
- 0.5% Chlorehexidine in 70% isopropyl alcohol
- · Products similar to optical cleaning liquid
- Bacillol AF
- Flux
- Sodium hypochlorite 10%

When selecting an alternative cleaning/disinfecting product, it is recommended to always identify the active ingredients. In case of doubt about a certain cleaning product, use plain water.

Do not use any of the following products:

- Alcohol in concentrations > 70%
- · Strong alkalis lye, strong solvents
- Acetone
- Toluene
- Acids
- Detergents containing fluoride
- · Detergents containing ammonia
- · Detergents containing abrasives
- Steel wool
- Sponge with abrasives
- · Steel blades
- · Cloths with steel thread
- · Paper-based cloths (e.g. paper towels, facial tissues, toilet paper)



CAUTION: Read and follow all instructions on the label of the cleaning product.



CAUTION: Take care not to damage or scratch the front glass or LCD. Be careful with rings or other jewelry and do not apply excessive pressure on the front glass or LCD.



CAUTION: When a small object or dust is tucked between the front bezel and the LCD surface (for displays without front glass), carefully remove with a soft object such as a plastic card or finger nail. Do not use sharp objects such as paperclips or tweezers to avoid damage to the LCD.



CAUTION: Do not apply or spray liquid directly to the display as excess liquid may cause damage to internal electronics. Instead, apply the liquid to a cleaning cloth.

Maintenance

Important information

6.1 Safety information

General recommendations

Read the safety and operating instructions before operating the device.

Retain safety and operating instructions for future reference.

Adhere to all warnings on the device and in the operating instructions manual.

Follow all instructions for operation and use.

Electrical Shock or Fire Hazard

To prevent electric shock or fire hazard, do not remove cover.

No serviceable parts inside. Refer servicing to qualified personnel.

Do not expose this apparatus to rain or moisture.

Modifications to the unit

Do not modify this equipment without authorization of the manufacturer.

Type of protection (electrical):

Display with external power supply: Class I equipment.

Degree of safety (flammable anesthetic mixture)

Equipment not suitable for use in the presence of a flammable anesthetic mixture with air or with oxygen or nitrous oxide.

Non-patient care equipment

- Equipment primarily for use in a health care facility that is intended for use where contact with a patient is unlikely (no applied part).
- The equipment shall not be used with life support equipment.
- The user should not touch the equipment, nor its signal input ports (SIP)/signal output ports (SOP) and the patient at the same time.

Power connection – Equipment with external 24 VDC power supply

- Power requirements: The equipment must be powered using the delivered medical approved 24 VDC (====) SELV power supply.
- The medical approved DC (====) power supply must be powered by the AC mains voltage.
- The power supply is specified as a part of the ME equipment or combination is specified as a ME system.
- To avoid the risk of electric shock, this equipment must only be connected to a supply mains with protective earth.
- · The equipment should be installed near an easily accessible outlet.
- The equipment is intended for continuous operation.

Transient over-voltage

If the device is not used for a long time, disconnect it from the AC inlet to avoid damage by transient over-voltage.

To fully disengage the power to the device, please disconnect the power cord from the AC inlet.

High magnetic environment

- The device shall not be used in the high magnetic environment of an MRI scanner.
- The installer shall assess the magnetic environment before installation or use of the device.

Power cords

Do not overload wall outlets and extension cords as this may result in fire or electric shock.

- Mains lead protection (U.S.: Power cord): Power cords should be routed so that they are not likely to be
 walked upon or pinched by items placed upon or against them, paying particular attention to cords at plugs
 and receptacles.
- Use a power cord that matches the voltage of the power outlet, which has been approved and complies with the safety standard of your particular country.
- Korea: Use KC certified products; Plug: 250 V~, 16 A; Power cord: 60227 IEC 53, 3G0.75 mm² / 60227 IEC 53, 3G1.0 mm²; Connector: 250 V~, 10 A

Grounding reliability

Grounding reliability can only be achieved when the equipment is connected to an equivalent receptacle marked "Hospital Only" or "Hospital Grade" (located on product or power supply cord)

Accessory equipment

Accessory equipment connected to the analog and digital interfaces must be in compliance with the respective nationally harmonized IEC standards (i.e. IEC 60950 for data processing equipment, IEC 60065 for video equipment, IEC 61010-1 for medical equipment). Furthermore, all configurations shall comply with the system standard IEC 60601-1-1. Anyone who connects additional equipment tot the signal input part or signal output part is configuring a medical system, and is therefore, responsible that the system complies with the requirements of the system standard IEC 60601-1-1. The unit is for exclusive interconnection with IEC 60601-1 certified equipment in the patient environment and IEC 60XXX certified equipment outside of the patient environment. If in doubt, consult the technical services department or your local representative.

Water and moisture

Never expose the display to rain or moisture.

Never use the display near water - e.g. near a bathtub, washbasin, swimming pool, kitchen sink, laundry tub or in a wet basement.

Ventilation

Do not cover or block any ventilation openings in the cover of the set. When installing the device in a cupboard or another enclosed location, heed the necessary space between the set and the sides of the cupboard.

Installation

Place the device on a flat, solid and stable surface that can support the weight of at least 3 devices. If you use an unstable cart or stand, the device may fall, causing serious injury to a child or adult, and serious damage to the device.

Malfunctions

Disconnect the equipment's power cord from the AC inlet and refer servicing to qualified service technicians under the following conditions:

- If the power cord or plug is damaged or fraved.
- If liquid has been spilled into the equipment.
- If the equipment has been exposed to rain or water.
- If the equipment does not operate normally when the operating instructions are followed. Adjust only those
 controls that are covered by the operating instructions since improper adjustment of other controls may
 result in damage and will often require extensive work by a qualified technician to restore the product to
 normal operation.
- If the equipment has been dropped or the cabinet has been damaged.
- If the product exhibits a distinct change in performance, indicating a need for service.

National Scandinavian Deviations for CL. 1.7.2

Finland: "Laite on liitettävä suojamaadoituskoskettimilla varustettuun pistorasiaan"

Norway: "Apparatet må tilkoples jordet stikkontakt" Sweden: "Apparaten skall anslutas till jordat uttag"

Operating principle

The single device output analog signals through ADC element (Analog Digital Convert) conversion to become a digital signal and the video signal is via Video Decoder conversion. It has become the same digital signal, these signals via Scaler IC as zoom in or out action and digital image processing, then through the cable line transmission LVDS signals to one of the LCD module. The last by the clock controller (Timing Controller, TCON), the clock signal is transmitted to the drive IC on the panel and turn on Backlight for LCD module light source by Scaler control.

Intended patient population: adults of infants.

BSMI

警語:使用過度恐傷害視力

注意事項:

- 1. 使用30分鐘請休息10分鐘
- 2. 未滿兩歲幼兒不看螢幕, 2歲以上每天看螢幕不要超過1小時

6.2 Cybersecurity

Hospital IT security

To prevent unauthorized access to the device, the organization incorporating the Nio 2MP LED Display in their IT network, shall have the necessary state-of-the-art policies, processes, standards and other security measures in place to incorporate, support and protect a medical device into the IT network. This shall include the application of risk management (e.g. by following IEC 80001-1:2010 or equivalent standards).

6.3 Environmental information

Disposal Information

Waste Electrical and Electronic Equipment



This symbol on the product indicates that, under the European Directive 2012/19/EU governing waste from electrical and electronic equipment, this product must not be disposed of with other municipal waste. Please dispose of your waste equipment by handing it over to a designated collection point for the recycling of waste electrical and electronic equipment. To prevent possible harm to the environment or human health from uncontrolled waste disposal, please separate these items from other types of waste and recycle them responsibly to promote the sustainable reuse of material resources.

For more information about recycling of this product, please contact your local city office or your municipal waste disposal service.

For details, please visit the Barco website at: http://www.barco.com/AboutBarco/weee

Turkey RoHS compliance



■ Türkiye Cumhuriyeti: AEEE Yönetmeliğine Uygundur.

[Republic of Turkey: In conformity with the WEEE Regulation]

中国大陆 RoHS

Chinese Mainland RoHS

根据中国大陆《电器电子产品有害物质限制使用管理办法》(也称为中国大陆RoHS),以下部分列出了 Barco产品中可能包含的有毒和/或有害物质的名称和含量。中国大陆RoHS指令包含在中国信息产业部MCV标 准:"电子信息产品中有毒物质的限量要求"中。 According to the "Management Methods for the Restriction of the Use of Hazardous Substances in Electrical and Electronic Products" (Also called RoHS of Chinese Mainland), the table below lists the names and contents of toxic and/or hazardous substances that Barco's product may contain. The RoHS of Chinese Mainland is included in the MCV standard of the Ministry of Information Industry of China, in the section "Limit Requirements of toxic substances in Electronic Information Products".

零件项目(名称)	有毒有害物质或元素						
Component name	Hazardous substances and elements						
	铅	汞	镉	六 价铬	多溴联苯	多溴二苯醚	
	Pb	Hg	Cd	Cr6+	PBB	PBDE	
印制电路配件	Х	0	0	0	0	0	
Printed Circuit Assemblies							
	X	0	0	0	0	0	
LCD panel							
外接电(线)缆	Х	0	0	0	0	0	
External Cables							
内部线路	0	0	0	0	0	0	
Internal wiring							
金属外壳	0	0	0	0	0	0	
Metal enclosure							
塑胶外壳	0	0	0	0	0	0	
Plastic enclosure							
散热片(器)	0	0	0	0	0	О	
Heatsinks							
风扇	0	0	0	0	0	О	
Fan							
电源供应器	Х	0	0	0	0	0	
Power Supply Unit							
文件说明书	0	0	0	0	0	0	
Paper Manuals							
光盘说明书	0	0	0	0	0	0	
CD manual 木丰校妹提S I/T 11364的坝宁绝线							

本表格依据SJ/T 11364的规定编制

This table is prepared in accordance with the provisions of SJ/T 11364.

- O: 表示该有毒有害物质在该部件所有均质材料中的含量均在 GB/T 26572 标准规定的限量要求以下.
- O: Indicates that this toxic or hazardous substance contained in all of the homogeneous materials for this part is below the limit requirement in GB/T 26572.
- X: 表示该有毒有害物质至少在该部件的某一均质材料中的含量超出 GB/T 26572 标准规定的限量要求.
- X: Indicates that this toxic or hazardous substance contained in at least one of the homogeneous materials used for this part is above the limit requirement in GB/T 26572.

在中国大陆销售的相应电子信息产品(EIP)都必须遵照中国大陆《电子电气产品有害物质限制使用标识要求》标准贴上环保使用期限(EFUP)标签。Barco产品所采用的EFUP标签(请参阅实例,徽标内部的编号使用于指定产品)基于中国大陆的《电子信息产品环保使用期限通则》标准。

All Electronic Information Products (EIP) that are sold within Chinese Mainland must comply with the "Marking for the restriction of the use of hazardous substances in electrical and electronic product" of Chinese Mainland, marked with the Environmental Friendly Use Period (EFUP) logo. The number inside the EFUP logo that Barco uses (please refer to the photo) is based on the "General guidelines of environment-friendly use period of electronic information products" of Chinese Mainland.



中国RoHS自我声明符合性标志 / China RoHS – SDoC mark

本产品符合《电器电子产品有害物质限制使用管理办法》和《电器电子产品有害物质限制使用达标管理目录》的要求。

This product meets the requirements of the "Management Rule on the Use Restriction of Hazardous Substances in Electrical and Electronic Products" and the "Management Catalogue for the Use Restriction of Hazardous Substances in Electrical and Electronic Products".



绿色自我声明符合性标志可参见电子档文件

The green SDoC mark is visible in the digital version of this document.

Taiwan BSMI RoHS

單元			限用	物質及其化學	符號	
Unit	Restricted substances and its chemical symbols			ls		
	鉛 Lead (Pb)	汞 Mercury (Hg)	鎘 Cadmium (Cd)	六價鉻 Hexavalent chromium (Cr ⁺⁶)	多溴聯苯 Polybrominat- ed biphenyls (PBB)	多溴二苯醚 Polybrominat- ed diphenyl ethers (PBDE)
印刷電路(含零组件) PCBA (including the components)	-	0	0	0	0	0
液晶顯示屏 LCD	-	0	0	0	0	0
電源适配器 Power adapter	-	0	0	0	0	0
電缆/連接器 Cable / connector	0	0	0	0	0	0
機械部件-金属件 Mechanical parts – Metal	-	0	0	0	0	0
塑膠殼 Plastic cover	0	0	0	0	0	0
電源線/信号線 Power supply / signal cable	-	0	0	0	0	0
機械部件-其他 Mechanical Parts – Others	0	0	0	0	0	0
紙/油墨(標簽·說明書·包裝盒等) Paper/Ink (Label, Instruction and packaging etc.)	0	0	0	0	0	0
背光燈管部件 Backlight assembly	0	0	0	0	0	0

備考1. "超出0.1 wt %"及 "超出0.01 wt %" 係指限用物質之百分比含量超出百分比含量基準值。

Note 1: "exceeding 0.1 wt%" and "exceeding 0.01 wt%" indicate that the percentage content of the restricted substance exceeds the reference percentage value of presence condition.

備考2. "O" 係指該項限用物質之百分比含量未超出百分比含量基準值。

Note 2: "O" indicates that the percentage content of the restricted substance does not exceed the percentage of reference value of presence.

備考3. "-" 係指該項限用物質為排除項目。

Note 3: The "-" indicates that the restricted substance corresponds to the exemption.

6.4 Regulatory compliance information

Indications for use

The display is intended to be used for displaying and viewing digital images (excluding digital mammography) for review and analysis by trained medical practitioners.

Intended usage environment

- The display is not in contact with patients.
- The display is not in the same environment as the patient.
- The display is intended to be used in a dedicated diagnostic reading room.

Caution (USA): Federal law restricts this device to sale by or on the order of a physician. (Details & exemptions are in the Code of Federal Regulations Title 21, 801 Part D).

Contra-indications

The device is not intended for digital mammography.

Intended users

Barco diagnostic displays are intended to be used for primary diagnosis by trained medical practitioners. The device is initially set up by trained integrators or medical IT staff.

Expected clinical benefits

The device aids in diagnosing patients.

Notice to the user and/or patient

Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.

Factory address

Coretronic Projection (Kunshan) Corp, No.20, 3rd Ave, Kunshan Free Trade Zone, Kunshan, Jiangsu, 215300 China

Manufacturing country

The manufacturing country of the product is indicated on the product label ("Made in ...").

Importers contact information

To find your local importer, contact one of Barco's regional offices via the contact information provided on our website (<u>www.barco.com</u>).

FCC class B

This device complies with Part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) this device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

This device has been tested and found to comply with the limits for a Class B digital device, pursuant to Part 15 of the FCC Rules. These limits are designed to provide reasonable protection against harmful interference in a residential installation. This device generates, uses and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation. If this device does cause harmful interference to radio or television reception, which can be determined by turning the device off and on, the user is encouraged to try to correct the interference by one or more of the following measures:

- Reorient or relocate the receiving antenna.
- Increase the separation between the device and receiver.
- Connect the device into an outlet on a circuit different from that to which the receiver is connected.

Consult the dealer or an experienced radio/TV technician for help.

Changes or modifications not expressly approved by the party responsible for compliance could void the user's authority to operate the equipment.

FCC responsible: Barco Inc., 3059 Premiere Parkway Suite 400, 30097 Duluth GA, United States, Tel: +1 678 475 8000

Canadian notice

CAN ICES-001(B) / NMB-001(B)

Brazilian local representative

BARCO LTDA, Av. Ibirapuera, 2332 - Torre II - 8° andar, Sao Paulo, Brazil

6.5 EMC notice

General information

This device is for use in professional healthcare facility environments only.

With the installation of the device, use only the delivered external cables and power supply or a spare part provided by the legal manufacturer. Using another can result in a decrease of the immunity level of the device.



WARNING: Use of this equipment adjacent to or stacked with other equipment should be avoided because it could result in improper operation. If such use is necessary, this equipment and the other equipment should be observed to verify that they are operating normally.



WARNING: Use of accessories, transducers and cables other than those specified or provided by the manufacturer of this equipment could result in increased electromagnetic emissions or decreased electromagnetic immunity of this equipment and result in improper operation.



WARNING: Portable RF communications equipment (including peripherals such as antenna cables and external antennas) should be used no closer than 30 cm (12 inches) to any part of the Nio 2MP LED Display, including cables specified by the manufacturer. Otherwise, degradation of the performance of this equipment could result.

Electromagnetic emissions

The Nio 2MP LED Display is intended for use in the electromagnetic environment specified below. The customer or the user of the Nio 2MP LED Display should assure that it is used in such an environment.

Emissions test	Compliance	Electromagnetic environment – Guidance
RF emissions CISPR 11	Group 1	The Nio 2MP LED Display uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emissions CISPR 11	Class B	The Nio 2MP LED Display is suitable for use in all
Harmonic emissions	Class D	establishments, including domestic establishments and those directly
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Complies	connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.

This Nio 2MP LED Display complies with appropriate medical EMC standards on emissions to, and interference from surrounding equipment. Operation is subject to the following two conditions: (1) this device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Interference can be determined by turning the equipment off and on.

If this equipment does cause harmful interference to, or suffer from harmful interference of, surrounding equipment, the user is encouraged to try to correct the interference by one or more of the following measures:

- Reorient or relocate the receiving antenna or equipment.
- Increase the separation between the equipment and receiver.
- Connect the equipment into an outlet on a circuit different from that to which the receiver is connected.
- Consult the dealer or an experienced technician for help.

Electromagnetic immunity

The Nio 2MP LED Display is intended for use in the electromagnetic environment specified below. The customer or the user of the Nio 2MP LED Display should assure that it is used in such an environment.

Immunity test	IEC 60601-1-2 test levels	Compliance level	Electromagnetic environment – guidance
Electrostatic discharge (ESD) IEC 61000-4-2	± 8 kV contact ± 2 kV, ± 4 kV, ± 8 kV, ± 15 kV air	± 8 kV contact ± 2 kV, ± 4 kV, ± 8 kV, ± 15 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%
Electrical fast transient/ burst IEC 61000-4-4	± 2 kV for power supply lines ± 1 kV for input/ output lines 100 kHz repetition frequency	± 2 kV for power supply lines ± 1 kV for input/ output lines 100 kHz repetition frequency	Mains power quality should be that of a typical commercial or hospital environment
Surge IEC61000-4-5	Line to line: ± 0.5 kV, ± 1 kV Line to ground: ± 0.5 kV, ± 1 kV, ± 2 kV	Line to line: ± 0.5 kV, ± 1 kV Line to ground: ± 0.5 kV, ± 1 kV, ± 2 kV	Mains power quality should be that of a typical commercial or hospital environment
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	period at 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315° 0% residual voltage for 1 period at 0° 70% residual voltage for 25 periods at 0°	0% residual voltage for 0.5 period at 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315° 0% residual voltage for 1 period at 0° 70% residual voltage for 25 periods at 0°	Mains power quality should be that of a typical commercial or hospital environment. If the user of the Nio 2MP LED Display requires continued operation during power mains interruptions, it is recommended that the Nio
	Voltage interruptions: 0% residual voltage for 250 periods at 0°	Voltage interruptions: 0% residual voltage for 250 periods at 0°	2MP LED Display be powered from an uninterruptible power supply or a battery
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	30 A/m	Not applicable ²	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment
Conducted RF IEC 61000-4-6	3 Vrms (6 Vrms in ISM bands) 150 kHz to 80 MHz	3 Vrms (6 Vrms in ISM bands)	-
Radiated RF IEC 61000-4-3	3 V/m 80 MHz to 2.7 GHz	3 V/m	

 $^{2. \}quad \hbox{Nio 2MP LED Display doesn't contain components that are susceptible to magnetic fields}.$

Immunity to RF wireless communications equipment

Test frequency (MHz)	Band (MHz)	Service	Modulation	Maximum power (W)	Distance (m)	Immunity test level (V/ m)	
385	380 – 390	TETRA 400	Pulse modulation 18 Hz	1.8	0.3	27	
450	430 – 470	GMRS 460, FRS 460	FM ± 5 kHz deviation 1 kHz sine	2	0.3	28	
710	704 – 787	LTE Band	Pulse	0.2	0.3	9	
745		13, 17	modulation 217 Hz				
780							
810	800 – 960	GSM 800/	Pulse	2	0.3	28	
870		900, TETRA 800, iDEN 820, CDMA 850, LTE Band 5	800, iDEN 820, CDMA 850, LTE	modulation 18 Hz			
930							
1720	1700 – 1990	GSM 1800,	Pulse	2	0.3	28	
1845		CDMA 1900, GSM 1900,	GSM 1900, GSM 1900,	modulation 217 Hz			
1970		DECT, LTE Band 1/3/4/ 25, UMTS					
2450	2400 – 2570	Bluetooth, WLAN, 802.11 b/g/n, RFID 2450, LTE Band 7	Pulse modulation 217 Hz	2	0.3	28	
5240	5100 – 5800	W LAN	Pulse	0.2	0.3	9	
5500		802.11 a/n	modulation 217 Hz				
5785							

6.6 Explanation of symbols

Symbols on the device

On the device or power supply, you may find the following symbols (nonrestrictive list):

((1639	Indicates the device meets the requirements of the applicable EC directives/ regulations.
F©	Indicates compliance with Part 15 of the FCC rules (Class A or Class B).
CI-NESSIFICE	Indicates the device is approved according to the UL regulations

Indicates the device is approved according to the UL regulations for Canada and US Indicates the device is approved according to the UL regulations for Canada and US Indicates the device is approved according to the UL Demko regulations. Indicates the device is approved according to the VCCI regulations. Indicates the device is approved according to the VCCI regulations. Indicates the device is approved according to the KC regulations. Indicates the device is approved according to the BSMI regulations. Indicates the device is approved according to the PSE regulations. Indicates the device is approved according to the RCM regulations. Indicates the device is approved according to the RCM regulations. Indicates the device is approved according to the EAC regulations. Indicates the device is approved according to the EAC regulations. Indicates the device is approved according to the BIS regulations. Indicates the device is approved according to the BIS regulations. Indicates the device is approved according to the BIS regulations. Indicates the device is approved according to the BIS regulations. Indicates the device is approved according to the INMETRO regulations.		
Indicates the device is approved according to the UL Demko regulations. Indicates the device is approved according to the CCC regulations. Indicates the device is approved according to the VCCI regulations. Indicates the device is approved according to the KC regulations. Indicates the device is approved according to the BSMI regulations. Indicates the device is approved according to the PSE regulations. Indicates the device is approved according to the RCM regulations. Indicates the device is approved according to the RCM regulations. Indicates the device is approved according to the EAC regulations. Caution: Federal law (United Stated of America) restricts this device to sale by or on the order of a licensed healthcare practitioner. Indicates the device is approved according to the BIS regulations. Indicates the device is approved according to the BIS regulations.	C UL US	Indicates the device is approved according to the UL regulations for Canada and US
Indicates the device is approved according to the VCCI regulations. Indicates the device is approved according to the VCCI regulations. Indicates the device is approved according to the KC regulations. Indicates the device is approved according to the BSMI regulations. Indicates the device is approved according to the PSE regulations. Indicates the device is approved according to the RCM regulations. Indicates the device is approved according to the EAC regulations. Caution: Federal law (United Stated of America) restricts this device to sale by or on the order of a licensed healthcare practitioner. Indicates the device is approved according to the BIS regulations. Indicates the device is approved according to the BIS regulations. Indicates the device is approved according to the BIS regulations.	CERTIFIED SAFETY US-CA E352529	Indicates the device is approved according to the UL regulations for Canada and US
Indicates the device is approved according to the KC regulations. Indicates the device is approved according to the KC regulations. Indicates the device is approved according to the BSMI regulations. Indicates the device is approved according to the PSE regulations. Indicates the device is approved according to the RCM regulations. Indicates the device is approved according to the EAC regulations. Indicates the device is approved according to the EAC regulations. Caution: Federal law (United Stated of America) restricts this device to sale by or on the order of a licensed healthcare practitioner. Indicates the device is approved according to the BIS regulations. Indicates the device is approved according to the BIS regulations.	D	Indicates the device is approved according to the UL Demko regulations.
Indicates the device is approved according to the KC regulations. Indicates the device is approved according to the BSMI regulations. Indicates the device is approved according to the PSE regulations. Indicates the device is approved according to the RCM regulations. Indicates the device is approved according to the EAC regulations. Caution: Federal law (United Stated of America) restricts this device to sale by or on the order of a licensed healthcare practitioner. Indicates the device is approved according to the BIS regulations. Indicates the device is approved according to the BIS regulations. Indicates the device is approved according to the BIS regulations.	(W)	Indicates the device is approved according to the CCC regulations.
Indicates the device is approved according to the BSMI regulations. Indicates the device is approved according to the PSE regulations. Indicates the device is approved according to the RCM regulations. Indicates the device is approved according to the EAC regulations. Caution: Federal law (United Stated of America) restricts this device to sale by or on the order of a licensed healthcare practitioner. Indicates the device is approved according to the BIS regulations. Indicates the device is approved according to the BIS regulations. Indicates the device is approved according to the BIS regulations.	[VEI]	Indicates the device is approved according to the VCCI regulations.
Indicates the device is approved according to the PSE regulations. Indicates the device is approved according to the RCM regulations. Indicates the device is approved according to the EAC regulations. Caution: Federal law (United Stated of America) restricts this device to sale by or on the order of a licensed healthcare practitioner. Indicates the device is approved according to the BIS regulations. Indicates the device is approved according to the INMETRO regulations.		Indicates the device is approved according to the KC regulations.
Indicates the device is approved according to the RCM regulations. Indicates the device is approved according to the EAC regulations. Caution: Federal law (United Stated of America) restricts this device to sale by or on the order of a licensed healthcare practitioner. Is 13232 (Part 1) IEC 08995-1 INDICE 08995-1 IND	3	Indicates the device is approved according to the BSMI regulations.
Indicates the device is approved according to the EAC regulations. Caution: Federal law (United Stated of America) restricts this device to sale by or on the order of a licensed healthcare practitioner. Is 13252 (Part 1)	PS E	Indicates the device is approved according to the PSE regulations.
Caution: Federal law (United Stated of America) restricts this device to sale by or on the order of a licensed healthcare practitioner. Is 13252 (Part 1)		Indicates the device is approved according to the RCM regulations.
the order of a licensed healthcare practitioner. Is 13252 (Part 1) Indicates the device is approved according to the BIS regulations. Indicates the device is approved according to the INMETRO regulations. Indicates the device is approved according to the INMETRO regulations.	ERE	Indicates the device is approved according to the EAC regulations.
R-XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX	$ m R_{ m only}$	
INMETRO	IEC 60950-1 R-XXXXXXXX	Indicates the device is approved according to the BIS regulations.
Indicates the USB connectors on the device.	INMETRO	Indicates the device is approved according to the INMETRO regulations.
	● ✓•	Indicates the USB connectors on the device.

Ð	Indicates the DisplayPort connectors on the device.
	Indicates the legal manufacturer.
	Indicates the manufacturing date.
	Indicates the entity importing the medical device into the locale.
xx J-yy	Indicates the temperature limitations ³ for the device to safely operate within specs.
MD	Indicates that the device is a Medical Device.
SN	Indicates the device Serial Number.
REF	Indicates the device part number or catalogue number.
UDI	Indicates the Unique Device Identifier.
EC REP	Indicates the Authorised Representative for the European Union.
CH REP	Indicates the Authorised Representative for Switzerland.
<u> </u>	Warning: dangerous voltage
	Caution
i	Consult the Instructions For Use.
elFU indicator	Consult the Instruction For Use on the website address that is provided as eIFU indicator.

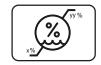
^{3.} Values for xx and yy can be found in the technical specifications paragraph.

	Indicates this device must not be thrown in the trash but must be recycled, according to the European WEEE (Waste Electrical and Electronic Equipment) directive.
	Indicates Direct Current (DC).
\sim	Indicates Alternating Current (AC).
(h)	Stand-by
\text{\$\rightarrow\$}\$	Equipotentiality
	Protective earth (ground)

Symbols on the box

On the box of the device, you may find the following symbols (nonrestrictive list):

T	Indicates a device that can be broken or damaged if not handled carefully when being stored.
*	Indicates a device that needs to be protected from moisture when being stored.
<u> </u>	Indicates the storage direction of the box. The box must be transported, handled and stored in such a way that the arrows always point upwards.
n or L	Indicates the maximum number of identical boxes which may be stacked on each other, where "n" is the limiting number.
xx-yy Kg or xx-yy Kg	Indicates the weight of the box and that it should be carried with two persons.
*	Indicates that the box should not be cut with a knife, a cutter or any other sharp object.
- xx*C	Indicates the temperature limits ⁴ to which the device can be safely exposed when being stored.



Indicates the range⁴ of humidity to which the device can be safely exposed when being stored.



Indicates the range⁴ of atmospheric pressure to which the device can be safely exposed when being stored.

6.7 Legal disclaimer

Disclaimer notice

Although every attempt has been made to achieve technical accuracy in this document, we assume no responsibility for errors that may be found. Our goal is to provide you with the most accurate and usable documentation possible; if you discover errors, please let us know.

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To protect our customers, Barco does not publicly disclose or confirm security vulnerabilities until Barco has conducted an analysis of the product and issued fixes and/or mitigations.

6.8 Technical specifications

Overview

Screen technology	IPS-Pro
Active screen size (diagonal)	540 mm (21.3")

Values for xx and yy can be found in the technical specifications paragraph.

Active screen size (H x V)	432 x 324 mm (17.0 x 12.8")
Aspect ratio (H:V)	4:3
Resolution	2MP (1600 x 1200 pixels)
Pixel pitch	0.27 mm
Color imaging	Yes
Gray imaging	Yes
Bit depth	30 bit
Viewing angle (H, V)	178°
Uniformity correction	ULT
Ambient light presets	Yes, reading room selection
Ambient light sensor	Yes
Backlight Output Stabilization (BLOS)	Yes
Front sensor	Yes
Maximum luminance (panel typical)	800 cd/m ²
DICOM calibrated luminance	500 cd/m ²
Contrast ratio (panel typical)	1400:1
Response time ((Tr + Tf)/2) (typical)	10 ms
Housing color	RAL 9003 / RAL 9004
Video input signals	1x DVI 1x DisplayPort
USB ports	1x USB 2.0 upstream (endpoint) 2x USB 2.0 downstream
Power rating	24 VDC, 3.75 A
Power requirements	This device shall only be powered by the following medical approved power supply: Bridgepower, type JMW190KB2400F11 Ratings marked on the medical power supply: Input rating: 100–240 VAC, 2 A, 50–60 Hz Output rating: 24 VDC, 3.75 A
Power consumption	50 W (nominal) < 1 W (hibernate) < 1 W (standby)
Dimensions with stand (W x H x D)	Portrait: 378 x 525~625 x 235 mm Landscape: 491 x 466~566 x 235 mm
Dimensions w/o stand (W x H x D)	Portrait: 378 x 491 x 83 mm Landscape: 491 x 378 x 83 mm
Dimensions packaged (W x H x D)	655 x 388 x 495 mm

Net weight with stand	With protective cover: 12.6 kg Without protective cover: 11.3 kg			
Net weight w/o stand	With protective cover: 7.6 kg Without protective cover: 6.3 kg			
Net weight packaged	With protective cover: 16.8 kg (without optional accessories) Without protective cover: 15.5 kg (without optional accessories)			
Tilt	-5° to +25°			
Swivel	-30° to +30°			
Pivot	90°			
Height adjustment range	100 mm			
Mounting standard	VESA (100 mm)			
Recommended modalities	All digital images, except digital mammography			
Certifications	CE1639 (MDD 93/42/EEC; A1:2007/47/EC class IIb product) IEC 60601-1(ed.3); IEC 60601-1(ed.3 am1) ANSI/AAMI ES60601-1(2005+C1:09+A2:10+A1:12) CAN/CSA-C22.2 No. 60601-1(2014) EN 60601-1-2:2015 EN 60601-1:2006/A11:2011/A12:2014; EN 60601-1:2006/A1:2013 CCC - GB9254-2008 + GB4943.1-2011 + GB17625.1-2012 KC BIS BSMI EAC INMETRO VCCI FCC class B ICES-001 Level B FDA 510(k) RoHS			
Supplied accessories	User guide Documentation disc System sheet Video cable Mains cable(s) USB cable External power supply			
QA software	QAWeb			
Warranty	5 years, including 20000 hrs backlight warranty			
Operating temperature	0 °C to 35 °C (15 °C to 30 °C within specs)			
Storage temperature	-20 °C to 60 °C			
Operating humidity	8% to 80% (non-condensing)			
Storage humidity	5% to 85% (non-condensing)			
L	ı			

Operating pressure	70 kPa minimum	
Storage pressure	50 to 106 kPa	

6.9 Open source license information

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 the distribution of this product by Barco.

Important information

Troubleshooting

7

7.1 Troubleshooting

General

If you experience trouble with the LCD display, refer to the following troubleshooting. If the problem persists, please contact your local dealer or our service center.

Problem: No image appears on the screen

- · Push the standby button.
- · Check if all I/O and power connectors are correctly connected as described in the "installation" section.
- Make sure the pins of the connectors are not crooked or broken.

Problem: Partial image or incorrectly displayed image

- Check to see if the resolution of your computer is higher than the resolution of the display.
- Re-configure the resolution of your computer to make it less than or equal to the native resolution (1600 x 1200)

7.2 Warning signals

General

Sometimes you may see warning messages for this display. This means that the display cannot correctly receive the signal form the computer graphics card.

Problem: No signal

This message means that the display has been powered on but it cannot receive any signal from the computer graphics card. Check all the power switches, power cables, and video cable.

Problem: Out of range

This message means that the signal of the computer graphics card is not compatible with the display. When the signal is not included in the compatibility mode we have listed in the appendices of this manual, the display will show this message.



