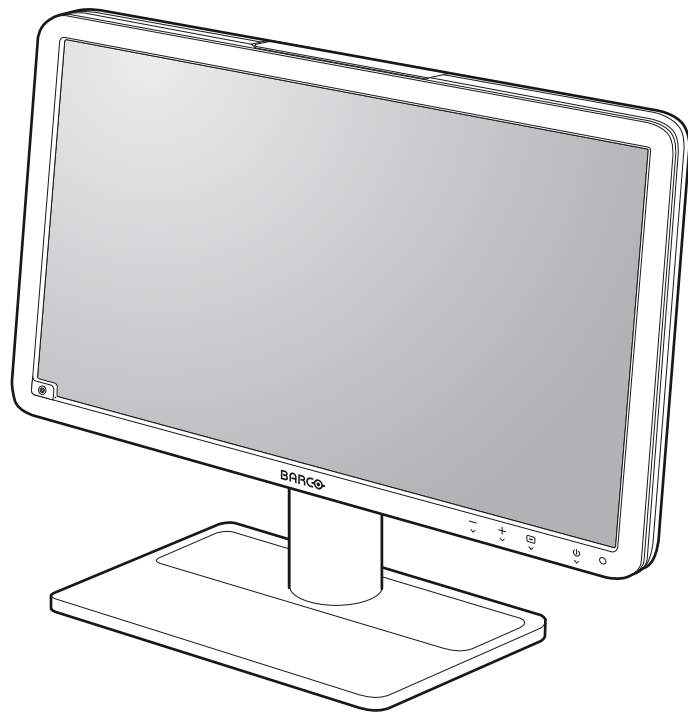# Eonis
22-inch clinical display
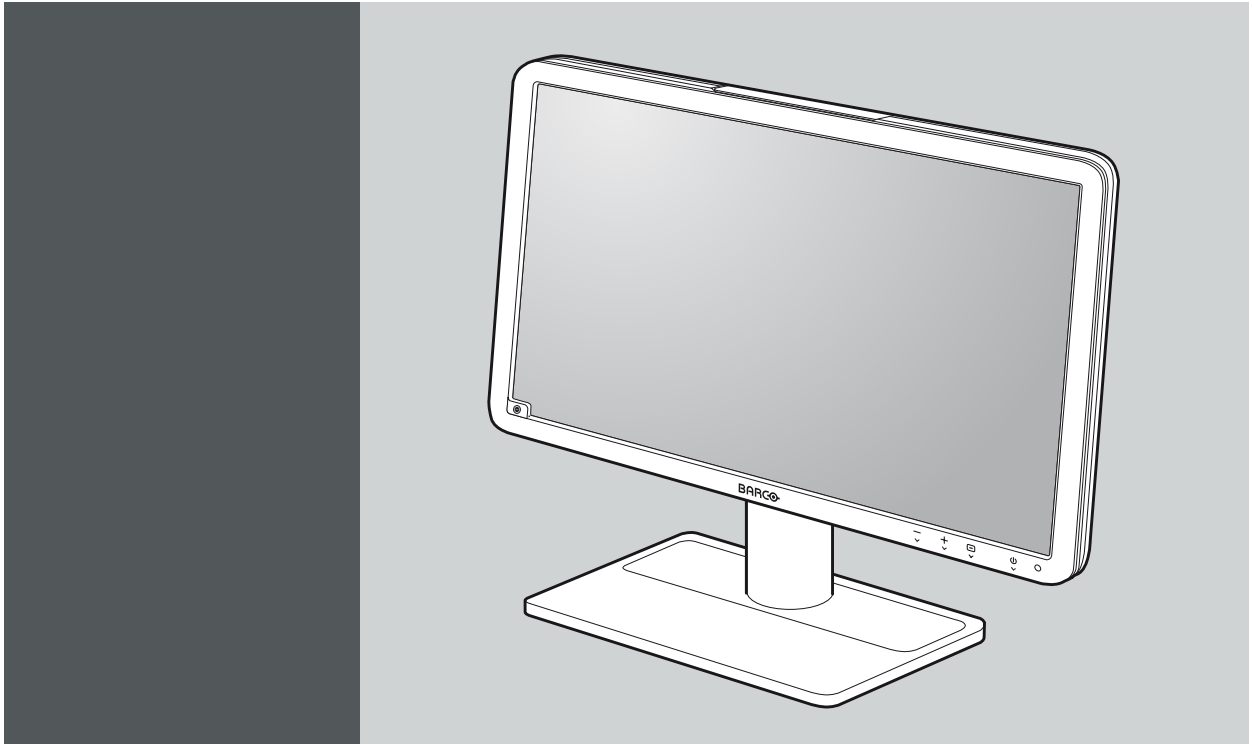
User Guide

MDRC-2122 WP
MDRC-2122 BL

BARCO

Visibly yours

Eonis

User Guide
22–inch clinical display

MDRC-2122 WP
MDRC-2122 BL

BARCO

Visibly yours

TABLE OF CONTENTS

1. WELCOME!

1.1 About the product

Consistent image quality

The Eonis display presents crisp, razor-sharp, high-contrast images. To guarantee image consistency at all times, it features a unique front sensor that automatically aligns image quality every time the display is turned on. This image consistency enhances collaboration: images always appear exactly like they are supposed to, on every display, which spurs confident clinical decisions. Furthermore, specialists looking at X-ray images will appreciate the built-in DICOM settings.

One-click quality assurance

Like Barco's entire medical display range, the Eonis display line comes with Barco's online MediCal QAWeb service for quality assurance and remote asset management. Proving its superior value in hospitals around the world, MediCal QAWeb provides automated quality assurance checks and detailed reports. It makes it easy for healthcare IT to centrally and remotely manage and configure displays across the healthcare organization. Also for private practices, QAWeb further strengthens image consistency and allows you to easily personalize your image.

Fully cleanable and smart design

The Eonis display carries all safety certifications required in clinical environments. It features an integrated cable management system, making it safe to use in hospitals and private practices. The multiple mounting options and connections further add to the smart design typical of Barco's Eonis display. Most unique, however, is the excellent cleanability of the white Eonis display thanks to the toughened and scratch-proof glass. This display model can be disinfected, both front and back, with typical alcohol-based cleaning agents.

Features

- Front sensor automatically aligns image quality
- IPS panel ensures wide viewing angle
- MediCal QAWeb suite for networked quality assurance, calibration and asset management all add to the exceptional, consistent image quality of the Eonis display
- Toughened, scratch-proof front glass (white version only)
- Covered cables
- Medical-grade certifications
- Sockets for Kensington lock makes the Eonis display safe for use in clinical environments
- Flexible VESA mount for easy arm, wall or cart mounting
- Top connections for easy and secure connections
- Multiple inputs (HDMI, DisplayPort, USB, etc.) ensure the display's flexible deployment

This manual further guides you through the different steps needed to install and use the Eonis display.

> ⚠️ **CAUTION: Read all the important safety information before installing and operating your Eonis display. Please refer to the dedicated chapter in this user guide.**

1.2 Symbols

Symbols in this document

The following symbols are used in this document:

⚠	Caution
⚡	Warning
ⓘ	Info, term definition. General info about the term
📄	Note: gives extra information about the described subject
👉	Tip: gives extra advice about the described subject

1.3 What's in the box

Overview

Your Eonis display comes with:

- this Eonis display user guide
- a documentation CD containing multilingual user documentation
- a system CD containing MediCal QAWeb Agent
- a (set of) AC power cord(s) (depending on the region of operation)
- a VGA cable
- a HDMI cable
- a USB cable
- an external power supply
- an accessory bag (cleaning cloth, velcro cable routing strap)

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

2. PARTS, CONTROLS AND CONNECTORS

2.1 Front view

Overview
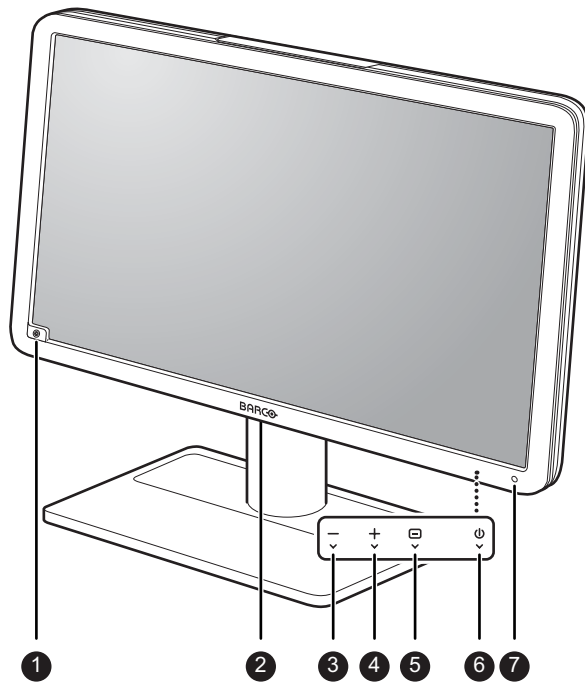
Image 2-1

1. Front sensor
2. USB downstream connector
3. Left / Decrease key
4. Right / Increase key
5. Menu / Enter key
6. Standby key
7. Power status LED

2.2 Rear view

Overview
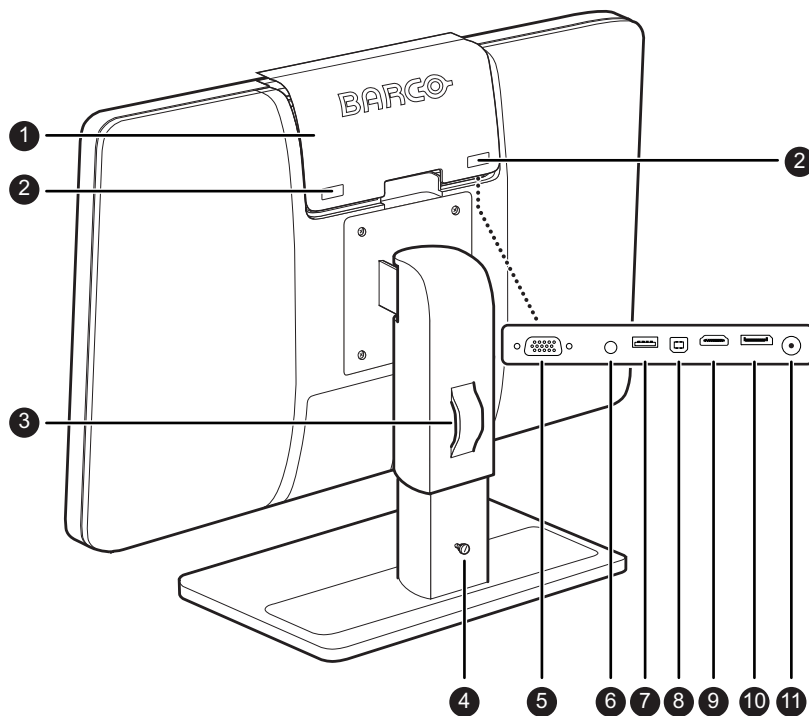
Image 2-2

1. Connector compartment cover
2. Kensington security slots
3. Opening for cable routing strap
4. Height mechanism lock screw
5. VGA input
6. HDMI audio line out
7. USB downstream connector
8. USB upstream connector
9. HDMI input
10. DisplayPort input
11. +12 VDC (⎓) power input

3. DISPLAY INSTALLATION

3.1 Unlocking the height mechanism

To unlock the height mechanism

In the factory, the height mechanism in the stand of your Eonis display is locked to prevent damage during transportation. You'll first have to unlock this mechanism before you can adjust your display height position.

To unlock the height mechanism, unscrew the height mechanism lock screw and remove it from the display stand.

Keep the lock screw in a known place for possible future transportation of your display.
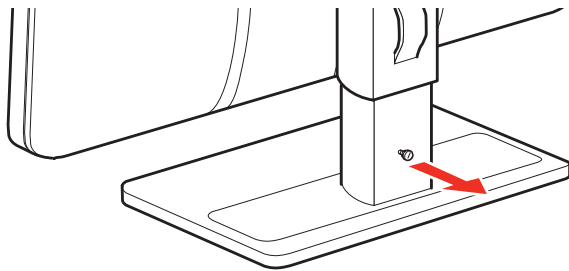
Image 3-1

3.2 Adjusting the display position

To adjust the display position

After unlocking the height mechanism, you can now safely tilt, swivel, raise and lower your display as desired.
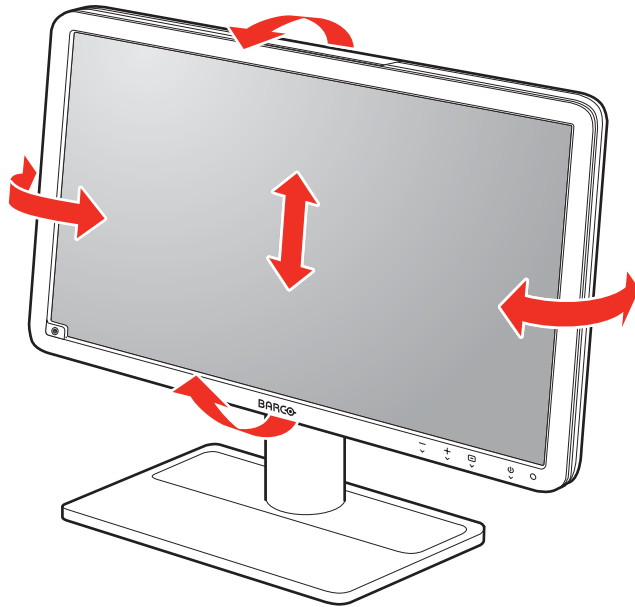
Image 3-2

⚠ **C**AUTION: **Do not try to pivot your display when attached to the stand. Trying to do so could cause serious damage to your display and its stand.**

3.3   Removing the cover

To remove the connector compartment cover

The connector compartment cover should be removed to get access to the connectors. To remove the cover, gently slide the connector compartment cover upwards and remove it from the display.
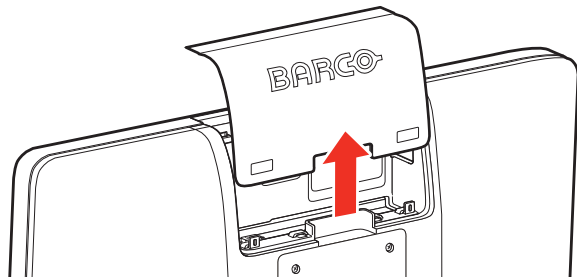
Image 3-3

3.4   Connecting the cables

To connect the cables

1. Connect one or more of the video input connections of your display (VGA, HDMI or DisplayPort) to the corresponding video outputs on your computer or any other video device.

2. If you want to make use of the display's USB downstream connectors, then connect your workstation with the display's USB upstream connector by means of the supplied USB cable.

3. Connect a speaker, amplifier, ... with the display's audio line out if you want to use the internal sound option of your display.

4. Connect the supplied external DC (━━) power supply to the +12 VDC power input of your display.

5. Plug the other end of the external DC (━━) power supply into a **grounded** power outlet by means of the proper power cord delivered with your display.
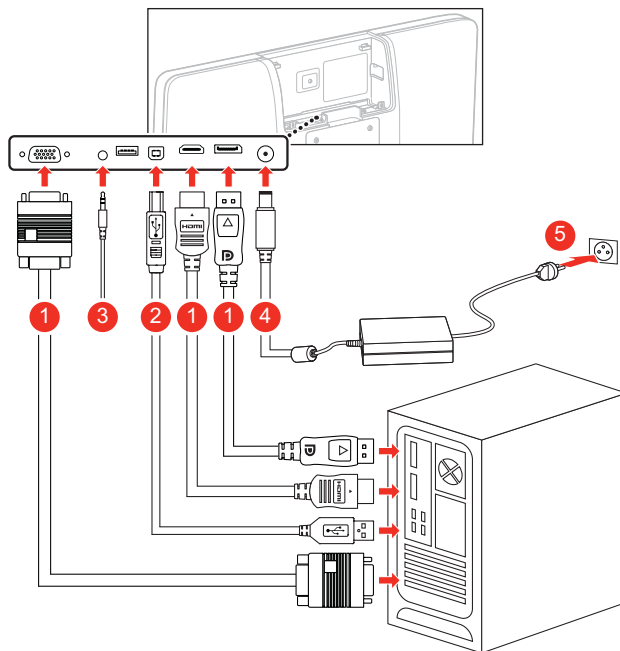
Image 3-4

3.5 Re-attaching the cover

To re-attach the connector compartment cover

Put the cover back in position by sliding it downwards so that it is fixed to the display again.

> **Pay attention that all cables stay in the cable channel while re-attaching the cover.**
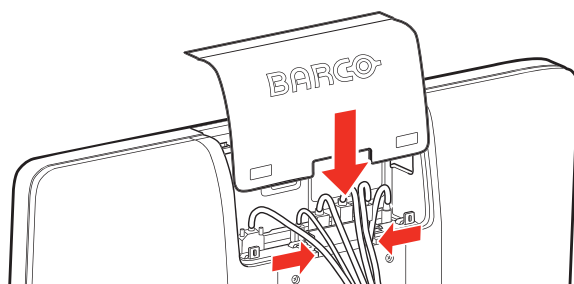
Image 3-5

3.6 Routing the cables

To route the cables

1. Slide the cable routing strap through the opening in the back of the stand.

2. Bundle all cables together so that they will fit in the strap.

3. Wrap and fix the cable routing strap around all cables.
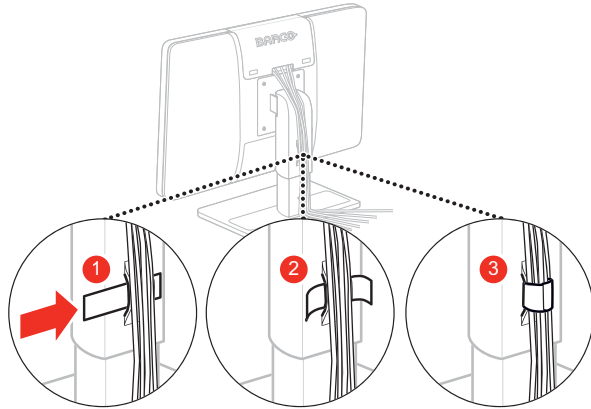
Image 3-6

3.7 Kensington security slots

To make use of the Kensington security slots

Your Eonis display has 2 Kensington slots available which allow you to secure the display to a desk or any other fixed object. Moreover, when locking the display with the connector compartment cover attached, you also prevent users from connecting/disconnecting any cables to/from the display.
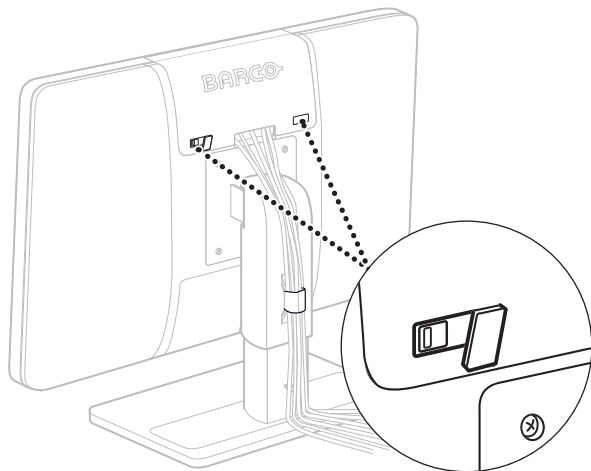
Image 3-7

3.8 VESA-mount installation

To mount the display on a VESA arm

The display panel, standard attached to a stand, is compatible with the VESA 75 mm standard.

1. Unscrew the four fixation screws to detach the panel from the stand.

2. Use 4 M4 screws to attach the panel to a VESA approved arm. Please make sure that the length of the screws is 10mm + VESA plate thickness (tolerance of +/- 1 mm).
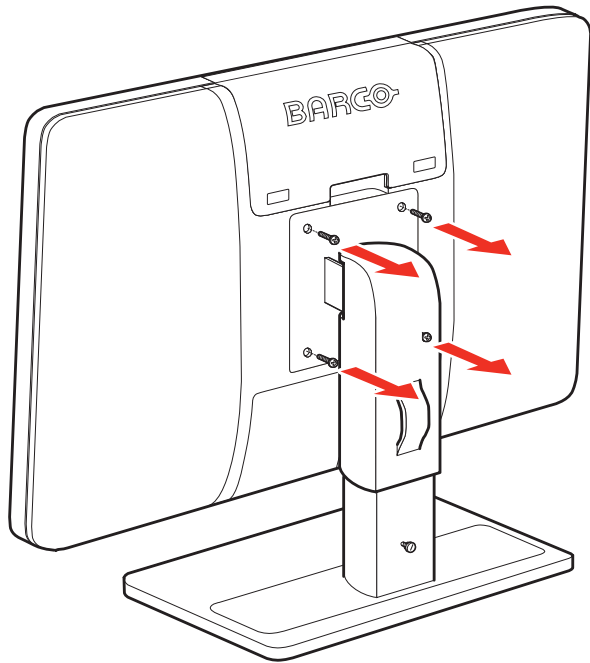
Image 3-8

> You should mount the panel in landscape position. Portrait position is possible but not supported.

> **CAUTION: Use an arm that can support a weight of at least 12 kg (26.50 lbs). Failure to do so could make the panel fall, causing serious injury to a child or adult, and serious damage to the equipment.**

> **CAUTION: 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 approved handle and use this to move the display. Please refer to the instruction manual of the arm for more information and instructions.**

4. DAILY OPERATION

4.1   Recommendations for daily operation

Optimize the lifetime of your display

Enabling the Display Power Management System (DPMS) of your display will optimize its diagnostic life-time 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 "Power Options Properties" in the "Control Panel".

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.

To learn more about LCD technology and missing pixels, consult the dedicated white papers available at www.barco.com/healthcare.

Maximize quality assurance

The 'MediCal QAWeb' system offers online service for high-grade Quality Assurance, providing maximum diagnostic confidence and uptime.

Barco recommends to install MediCal QAWeb Agent and apply the default QAWeb policy at least. This policy includes calibration on regular intervals. Upgrading to MediCal QAWeb Server offers even more possibilities for hospitals. Learn more and sign up for the free MediCal QAWeb Essential level at www.barco.com/healthcare/qa

4.2   On/Off switching

To switch your display on or off

1. Shortly press the Standby (⏻) key.

4.3   Bringing up the OSD menus

About the OSD menu

The OSD menu allows you to configure different settings to make your Eonis 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 bring up the OSD menu

1. While the display is switched on, press the Menu/Enter (▣) key.
   As a result, the OSD main menu comes up in the middle of the screen. If no further actions are taken within the following 10 seconds however, the OSD menu will disappear again.

4.4   Navigating through the OSD menus

To navigate through the OSD menus

1. Use the Right/Down (+) and Left/Up (−) keys to move through the (sub)menus, change values or make selections.

2. To go into a submenu or confirm adjustments and selections, use the Menu/Enter (▣) key.

5. ADVANCED OPERATION

5.1   Video input source selection

About video input source selection

By default, your Eonis display automatically detects and shows the connected video input source. However, when for instance more then one video input source is connected, it may be needed to manually select the input source to be displayed.

The available video input source selections for your display are:

* *Auto*: This is the default setting and will automatically detect and display the connected video input source.
* *HDMI*: This setting will display the video connected to the HDMI input.
* *DisplayPort*: This setting will display the video connected to the DisplayPort input.
* *VGA*: This setting will display the video connected to the VGA input.

To select a video input source

1. Bring up the OSD main menu.

2. Navigate to the *Input Selection* menu.

3. Select one of the available video input sources.

5.2   Luminance adjustment

About luminance adjustment

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

To adjust the luminance

1. Bring up the OSD main menu.

2. Navigate to the *Adjustments* menu.

3. Enter the *Luminance* submenu.

4. Set a luminance value as desired and confirm.

5.3   Gamma selection

About gamma selection

Native, uncorrected panels will display all grayscale/color levels with equal luminance increments. Studies have shown however, that in medical images certain grayscale/color parts contain more diagnostic information then others. To respond to these conclusions, gamma functions have been defined. These functions emphasize on these parts containing crucial diagnostic information by correcting the native panel behavior.

The available gamma functions for your display are:

- *Native*: If you select *Native*, the native panel behavior will not be corrected.
- *sRGB*: This is the display function as defined in the sRGB specification and is designed to match typical home and office viewing conditions. It is widely used in most computer applications.
- *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 gamma function results in more visible grayscales in the images. Barco recommends selecting the *DICOM* gamma function for most medical viewing applications.
- *Gamma 2.2*: Select this function in case the display is to replace a CRT display with a gamma of 2.2.
- *QAWeb*: This gamma function will be automatically selected when gamma functions are defined by MediCal QAWeb.

To select a gamma function

1. Bring up the OSD main menu.

2. Navigate to the *Adjustments* menu.

3. Enter the *Gamma* submenu.

4. Select one of the available gamma functions.

5.4   Ambient light reading room selection

About ambient light reading rooms

The available ambient light reading rooms for your display are:

- *Dark Room*: Corresponds to light conditions in dark diagnostic reading rooms. This setting has the lowest maximum ambient light.
- *Office*: Corresponds to light conditions in office rooms.
- *Operation Room*: Corresponds to light conditions in operating rooms. This setting has the highest maximum ambient light.
- *QAWeb*: This setting will be automatically selected when ambient light conditions are defined by MediCal QAWeb.

To select an ambient light reading room

1. Bring up the OSD main menu.

2. Navigate to the *Adjustments* menu.

3. Enter the *Ambient Light* submenu.

4. Select one of the available reading rooms and confirm.

5.5   White point selection

About white point selection

This setting allows you to modify the display white point, used as reference for all other colors to be displayed.

The available white point settings for your display are:

- *Native*: The native, unmodified color temperature of the LCD panel.
- *6500K (sRGB)*: Corresponds to a color temperature of 6500 Kelvin (D65).
- *QAWeb*: This white point setting will be automatically selected when white point is defined by MediCal QAWeb.

To select the white point

1. Bring up the OSD main menu.

2. Navigate to the *Adjustments* menu.

3. Enter the *White Point* submenu.

4. Select one of the available white point presets.

5.6   Analog video settings

> **The following settings are only available when VGA video input source is selected.**

About analog video settings

When the VGA video input source is active, a number of analog video settings will become available:

• *Auto Adjust*: The analog video setting will automatically be adjusted
• *Geometry*: Allows to manually adjust the geometry settings of the analog video (clock frequency, clock phase, horizontal position, vertical position)
• *Level*: Allows to manually adjust the contrast and brightness levels of the analog video

To adjust the analog video settings

1. Bring up the OSD main menu.

2. Navigate to the *Adjustments* menu.

3. Enter the *Analog* submenu.

4. Adjust one of the available analog video settings as desired.

5.7   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 Eonis display:

• English
• French
• German
• Spanish
• Italian
• Dutch
• Japanese
• Traditional Chinese
• Simplified Chinese
• Korean

To select the language of the OSD menu:

1. Bring up the OSD main menu.

2. Navigate to the *Adjustments > Settings* menu.

3. Enter the *Language* submenu.

4. Select one of the available languages.

5.8   Power status LED

About the power status LED

By default, the power status LED has the following behavior:

- **Green:** Display is on
- **Blinking green:** Display is entering standby power-saving mode
- **Orange:** Display is in standby power-saving mode
- **Off:** Display is disconnected from the mains power

This default behavior can be changed so that the power status LED is also off when the display is on or when the display is entering standby power-saving mode.

> **The orange standby power-saving state of the LED is not influenced by this setting. So, when the display is in standby power-saving mode, the LED will turn orange, even if it was switched off by this setting.**

To change the behavior of the power status LED:

1. Bring up the OSD main menu.

2. Navigate to the *Adjustments > Settings* menu.

3. Enter the *Power Status LED* submenu.

4. Change the behavior of the power status LED as desired and confirm.

5.9   DPMS mode

About DPMS mode

Enabling the Display Power Management System (DPMS) mode on your Eonis display will optimize the displays' lifetime by automatically switching off the backlight when no video signal is detected for approximately 10 seconds. The power status LED will then turn orange.

To enable/disable DPMS mode

1. Bring up the OSD main menu.

2. Navigate to the *Adjustments > Settings > Power Save* menu.

3. Enter the *DPMS* submenu.

4. Select *On* or *Off* as desired and confirm.

5.10 Proximity sensor time out

About the proximity sensor

As an optional accessory, a Barco proximity sensor can be connected to your Eonis display which will detect if somebody is present in front of the display. To optimize the displays' lifetime, the backlight can automatically be switched off after a specified time of inactivity (i.e. when nobody is present). As soon as the proximity sensor detects that somebody is present again, the backlight will automatically be switched on again. The available time-out values for this power saving mechanism are:

- *30 sec*
- *2 min*
- *10 min*
- *Off*

> **Please note that the this power saving mechanism will only work when also DPMS mode is enabled.**

To adjust the time out of the proximity sensor

1. Bring up the OSD main menu.

2. Navigate to the *Adjustments > Settings > Power Save* menu.

3. Enter the *Optional Proximity Sensor* submenu.

4. Select one of the available time out presets.

5.11 Self calibration frequency

About self calibration

The front sensor of your Eonis display measures the output luminance of your screen and allows the display to automatically stabilize its luminance for maximum image quality over the displays' lifetime. This self calibration is done at an adjustable, predefined frequency:

- *1 min*
- *1 hr*
- *6 hr*
- *24 hr*
- *Never*
- *QAWeb*: This setting will be automatically selected when the self calibration frequency is defined by MediCal QAWeb.

To adjust the self calibration frequency

1. Bring up the OSD main menu.

2. Navigate to the *Adjustments > Settings* menu.

3. Enter the *Self Calibration Frequency* submenu.

4. Select one of the available frequency presets.

5.12 Factory reset

About factory reset

A factory reset allows you to fully restore the display to its original factory setting.

To perform a factory reset

1. Bring up the OSD main menu.

2. Navigate to the *Adjustments > Settings* menu.

3. Enter the *Factory Reset* submenu.

4. Select *Yes* or *No* as desired and confirm.

6. CLEANING YOUR DISPLAY

6.1   Cleaning instructions

To clean the display

Clean the display using a sponge, cleaning cloth or soft tissue, lightly moistened with a recognized cleaning product for medical equipment. Read and follow all label instructions on the cleaning product. In case of doubt about a certain cleaning product, use plain water.

Do not use following products:

- Alcohol/solvents at higher concentration > 5%
- Strong alkalis lye, strong solvents
- Acid
- Detergents with fluoride
- Detergents with ammonia
- Detergents with abrasives
- Steel wool
- Sponge with abrasives
- Steel blades
- Cloth with steel thread

⚠ **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: For model MDRC-2122 BL only: Do not apply or spray liquid directly to the display as excess liquid may cause damage to internal electronics. Instead, apply the liquid to the cleaning cloth.**

7. IMPORTANT INFORMATION

7.1   Safety information
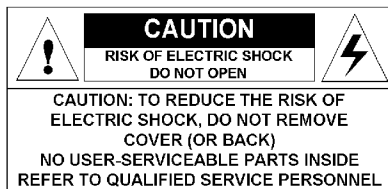
General recommendations

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

```
        ⚠                CAUTION                ⚡
                  RISK OF ELECTRIC SHOCK
                        DO NOT OPEN
         CAUTION: TO REDUCE THE RISK OF
         ELECTRIC SHOCK, DO NOT REMOVE
                   COVER (OR BACK)
         NO USER-SERVICEABLE PARTS INSIDE
       REFER TO QUALIFIED SERVICE PERSONNEL
```

• Do not modify this equipment without authorization of the manufacturer.
• No user-serviceable part inside. The equipment should be opened only by qualified service personnel.

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 may not be used with life support equipment.
• The user should not touch the equipment and the patient at the same time.

Power connection – Equipment with external 12 VDC power supply

• Power requirements: The equipment must be powered using the delivered medical approved 12 VDC ($=\!=\!=$) 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.

Power cords:

- Utilize a UL-listed detachable power cord, 3-wire, type SJ or equivalent, 18 AWG min., rated 250 V min., provided with a hospital-grade type plug 5-15P configuration for 120V application, or 6-15P for 240V application.
- 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.
- The power supply cord should be replaced by the designated operator only at all time.

External equipment

External equipment intended for connection to signal input/output or other connectors, shall comply with relevant UL/ EN/ IEC standard (e.g. UL/EN/IEC 60950 for IT equipment and UL/EN 60601-1 / IEC 60601 series for medical electrical equipment). In addition, all such combinations -systems- shall comply with the standard IEC 60601-1-1, safety requirements for medical electrical systems. Equipment not complying with UL/EN / IEC 60601-1 shall be kept outside the patient environment, as defined in the standard.

Equipment not complying with IEC 60601 must be kept outside the patient environment, as defined in the standard as at least 1.5 meters from the patient or the patient support.

Any person who connects external equipment to signal input, signal output, or other connectors has formed a system and is therefore responsible for the system to comply with the requirements of IEC 60601-1-1. If in doubt, speak with a qualified technician.

In locations where 240 V outlets are used, connect this display only on a center-tapped, 240 V, single-phase supply.

Water and moisture

Never expose the device to rain or moisture. Never use the device near water - e.g. near a bathtub, washbasin, swimming pool, kitchen sink, laundry tub or in a wet basement.

IP-x level for MDRC-2122 BL: Ordinary

IP-x level for MDRC-2122 WP: Protection glass

Ventilation

Do not cover or block any ventilation openings in the cover of the set. When installing the device in a cupboard or another closed 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.

This apparatus conforms to:

CE (MDD 93/42/EEC class I product), CE - 2004/ 108/EC, IEC 60950-1:2005 + A1:2009 (2ND EDITION), IEC 60601-1 2ND ED:1988 + A1:1991 + A2:1995, UL 60601-1 1ST EDITION, CAN/CSA-C22.2 NO. 601.1-M90:2005, IEC 60601-1:2005 + A1:2012, ANSI/AAMI ES 60601-1:2005 + C1:2009 + A1:2012, CAN/CSAC22.2 No. 60601-1-08:2008, DEMKO - EN 60601- 1:2006, EN 60601-1-2:2007, CCC - GB9254-2008 + GB4943-2001 + GB17625.1-2003, KC, VCCI, FCC class B, ICES-001 Level B, FDA Class I device, RoHS

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"

7.2   Environmental information

Disposal information (Waste Electrical and Electronic Equipment)

This symbol on the device indicates that, under European Directive 2002/96/EC governing waste from electrical and electronic equipment, this device must not be disposed of with other municipal waste. Please dispose of your waste device 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 devices from other types of waste and recycle them responsibly to promote the sustainable reuse of material resources.

For more information about recycling of this device, please contact your local city office or your municipal waste disposal service. For details, please visit the Barco website at: http://www.barco.com/en/About-Barco/weee.

Turkey RoHS compliance

Republic of Turkey: In conformity with the EEE Regulation

Türkiye Cumhuriyeti: EEE Yönetmeliğine Uygundur

中国大陆 RoHS

Chinese Mainland RoHS

根据中国大陆《电子信息产品污染控制管理办法》(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 "China Administration on Control of Pollution Caused by Electronic Information 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) Component name	有毒有害物质或元素 Hazardous substances and elements					
	铅 Pb	汞 Hg	镉 Cd	六价铬 Cr6+	多溴联苯 PBB	多溴二苯醚 PBDE
印制电路配件 Printed Circuit Assemblies	X	O	O	O	O	O
液晶面板 LCD panel	X	O	O	O	O	O
外接电(线)缆 External Cables	X	O	O	O	O	O
內部线路 Internal wiring	O	O	O	O	O	O

零件项目(名称)	有毒有害物质或元素					
Component name	**Hazardous substances and elements**					
	铅 **Pb**	汞 **Hg**	镉 **Cd**	六价铬 **Cr6+**	多溴联苯 **PBB**	多溴二苯醚 **PBDE**
金属外壳 Metal enclosure	O	O	O	O	O	O
塑胶外壳 Plastic enclosure	O	O	O	O	O	O
散热片(器) Heatsinks	O	O	O	O	O	O
电源供应器 Power Supply Unit	X	O	O	O	O	O
风扇 Fan	O	O	O	O	O	O
文件说明书 Paper Manuals	O	O	O	O	O	O
光盘说明书 CD manual	O	O	O	O	O	O
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 SJ/T 11363-2006 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 SJ/T11363-2006.						
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 SJ/T 11363-2006 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 SJ/T11363-2006						

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《电子信息产品污染控制标识要求》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制定产品）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 "Electronic Information Products Pollution Control Labeling Standard" 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 "Standard of Electronic Information Products Environmental Friendly Use Period" of Chinese Mainland.

7.3 Regulatory compliance information

Indications for use

This display is an AMLCD display designed for viewing medical X-ray images. This unit should not be used near patients (where patients are likely to be in unconscious condition) and should be kept outside of 1.83 m perimeter and 2.29 m vertical.

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.

Canadian notice

This ISM device complies with Canadian ICES-001.

Cet appareil ISM est conforme à la norme NMB-001 du Canada.

7.4   EMC notice

General information

No specific requirement on the use of external cables or other accessories except power supply.

With the installation of the device, use only the delivered 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.

Electromagnetic emissions

The Eonis display is intended for use in the electromagnetic environment specified below. The customer or the user of the Eonis display should assure that it is used in such an environment.

Emissions test	Compliance	Electromagnetic environment – Guidance
RF emissions CISPR 11	Group 1	The Eonis 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 Eonis display is suitable for use in all establishments, including domestic establishments and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.
Harmonic emissions IEC 61000-3-2	Class D	
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Complies	

This Eonis 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 Eonis display is intended for use in the electromagnetic environment specified below. The customer or the user of the Eonis display should assure that it is used in such an environment.

Immunity test	IEC 60601 Test levels	Compliance level	Electromagnetic environment – guidance
Electrostatic discharge (ESD) IEC 61000-4-2	± 6kV contact ± 8kV air	± 6kV contact ± 8kV 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	± 2kV for power supply lines ± 1kV for input/ output lines	± 2kV for power supply lines ± 1kV for input/ output lines	Mains power quality should be that of a typical commercial or hospital environment
Surge IEC61000-4-5	± 1 kV line(s) to line(s) ± 2 kV line(s) to earth	± 1 kV line(s) to line(s) ± 2 kV line(s) to earth	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	< 5% U_T [1] (> 95% dip in U_T) for 0.5 cycle 40% U_T (60% dip in U_T) for 5 cycles 70% U_T (30% dip in U_T) for 25 cycles < 5% U_T (>95% dip in U_T) for 5s	< 5% U_T (> 95% dip in U_T) for 0.5 cycle 40% U_T (60% dip in U_T) for 5 cycles 70% U_T (30% dip in U_T) for 25 cycles < 5% U_T (>95% dip in U_T) for 5s	Mains power quality should by that of a typical commercial or hospital environment. If the user of the Eonis display requires continued operation during power mains interruptions, it is recommended that the Eonis display be powered from an uninterruptible power supply or a battery.
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	3 A/m	Not applicable	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 Radiated RF IEC 61000-4-3	3 Vrms 150 kHz to 80 MHz 3 V/m 80 MHz to 2.5 GHz	3 V 3 V/ m	Portable and mobile RF communications equipment should be used no closer to any part of the Eonis display, including cables, than the recommended

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1. is the a.c. mains voltage prior to application of the test level.

Immunity test	IEC 60601 Test levels	Compliance level	Electromagnetic environment – guidance
			separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance **d = 1.2√P** **d = 1.2√P** 80 MHz to 800 MHz **d = 2.3√P** 800 MHz to 2.5 Ghz Where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in meters (m). Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey,[2] should be less than the compliance level in each frequency range.[3] Interference may occur in the vicinity of equipment marked with symbol:

At 80 MHz and 800 MHz, the higher frequency range applies.

These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

2. Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the Eonis display is used exceeds the applicable RF compliance level above, the Eonis display should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the Eonis display.
3. Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.

Recommended separation distance

The Eonis display is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer of the user of the Eonis display can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the Eonis display as recommended below, according to the maximum output power of the communications equipment.

Rated maximum output power of transmitter [4] W	Separation distance according to frequency of transmitter		
	150kHz to 80MHz $d=1.2\sqrt{P}$	80MHz to 800MHz $d=1.2\sqrt{P}$	800MHz to 2.5GHz $d=2.3\sqrt{P}$
0.01	0.12	0.12	0.23
0.1	0.38	0.38	0.73
1	1.2	1.2	2.3
10	3.8	3.8	7.3
100	12	12	23

At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.

These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection form structures, object and people.
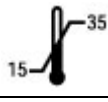
7.5   Explanation of symbols

Symbols on the device

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

CE	Indicates compliance with the Directive 93/42/EEC as Class I device
CE 0120	Indicates compliance with the Directive 93/42/EEC as Class II device
FC	Indicates compliance with Part 15 of the FCC rules (Class A or Class B)
CLASSIFIED UL	Indicates the device is approved according to the UL regulations

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
4. For transmitters rated at a maximum output power not listed above, the recommended separation distance d in meters (m) can be estimated using the equation applicable to the frequency of the transmitter. Where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer.

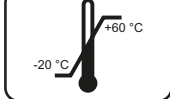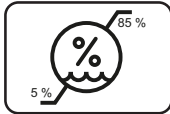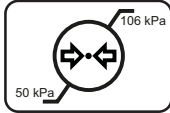|	
	Indicates the device is approved according to the UL regulations for Canada and US
	Indicates the device is approved according to the UL Demko regulations
	Indicates the device is approved according to the CCC regulations
	Indicates the device is approved according to the VCCI regulations
	Indicates the device is approved according to the KC regulations
	Indicates the device is approved according to the BSMI regulations
	Indicates the USB connectors on the device
	Indicates the DisplayPort connectors on the device
	Indicates the manufacturing date
	Indicates the temperature limitations for the device to safely operate within specs
	Indicates the device serial no.
	Consult the operating instructions
	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)
	Indicates Alternating Current (AC)

Symbols on the box

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

	Indicates a medical device that can be broken or damaged if not handled carefully when being stored.
	Indicates a medical device that needs to be protected from moisture when being stored.
	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.
15	Indicates the maximum number of boxes to be stacked on each other.
20 - 30 Kg	Indicates that the box should be carried with two persons.
	Indicates that the box should not be cut with a knife, a cutter or any other sharp object.
+60 °C -20 °C	Indicates the temperature limits to which the medical device can be safely exposed when being stored.
85 % 5 %	Indicates the range of humidity to which the medical device can be safely exposed when being stored.
106 kPa 50 kPa	Indicates the range of atmospheric pressure to which the medical device can be safely exposed when being stored.

7.6   **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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7.7   Technical specifications

MDRC-2122 BL

Product acronym	MDRC-2122 BL
Screen technology	TFT Color LCD
Active screen size (diagonal)	542 mm (21.5")
Active screen size (H x V)	267,3 mm x 475,2 mm (10.52" x 18.71")
Aspect ratio (H:V)	16:9
Resolution	2MP (1920 x 1080)
Pixel pitch	0.2475
Color imaging	Yes
Gray imaging	Yes
Color support	10 bit; 16,7 million colors
Viewing angle (H, V)	178°
Ambient Light Compensation (ALC)	Yes, preset values in OSD
Front sensor	Yes
Maximum luminance	250 cd/m² typical
DICOM calibrated luminance	180 cd/m²
Contrast ratio	1000:1 typical
Response time (Tr + Tf)	14 ms
Housing color	Black + Silver
Video input signals	VGA, DisplayPort, HDMI
USB ports	1 upstream, 2 downstream
USB standard	2.0
Power requirements (nominal)	100-240 V
Power consumption (nominal)	40 W (<0,5 W in stand-by)
Power save mode	Yes
Power management	DPMS
Dot clock	165 MHz
OSD languages	English, French, German, Spanish, Italian, Dutch, Japanese, Traditional Chinese, Simplified Chinese, Korean
Dimensions with stand (W x H x D)	507 x 377–457x 166 mm
Dimensions w/o stand (W x H x D)	507 x 306 x 57 mm
Dimensions packaged (W x H x D)	610 x 505 x 230 mm
Net weight with stand	6.0 kg
Net weight w/o stand	3.6 kg
Net weight packaged with stand	9.6 kg
Height adjustment range	80 mm
Tilt	0° / +20°
Swivel	-30° / +30°
Pivot	N/A
Mounting standard	VESA (75 mm)
Screen protection	N/A

Certifications	CE (MDD 93/42/EEC class I product), CE - 2004/ 108/EC, IEC 60950-1:2005 + A1:2009 (2ND EDITION), IEC 60601-1 2ND ED:1988 + A1:1991 + A2:1995, UL 60601-1 1ST EDITION, CAN/CSA-C22.2 NO. 601.1-M90:2005, IEC 60601-1:2005 + A1:2012, ANSI/AAMI ES 60601-1:2005 + C1:2009 + A1:2012, CAN/CSAC22.2 No. 60601-1-08:2008, DEMKO - EN 60601- 1:2006, EN 60601-1-2:2007, CCC - GB9254-2008 + GB4943-2001 + GB17625.1-2003, KC, VCCI, FCC class B, ICES-001 Level B, FDA Class I device, RoHS
Supplied accessories	User Guide
	Cable routing strap
	Video cables (1 x VGA + 1 x HDMI)
	Main cables (UK, European (CEBEC/KEMA) or USA (UL/ CSA; adaptor plug NEMA 5-15P) or Chinese (CCC))
	USB 2.0 cable
	External power supply (model: BridgePower Corp. BPM060S12F15; Input: 100-240 V AC, 50-60 Hz, 1.5 A; Output: +12V DC (⎓), 5.0 A)
Optional accessories	Proximity sensor
QA software	MediCal QAWeb 2
Warranty	3 years
Operating temperature	10°C - 40°C
Storage temperature	-20°C - 60°C
Operating humidity	30% - 75% (non-condensing)
Storage humidity	5% - 85% (non-condensing)
Operation altitude	2000 m
Storage altitude	7500 m
Operating pressure	70 kPa - 106 kPa
Storage pressure	50 kPa - 106 kPa

MDRC-2122 WP

Product acronym	MDRC-2122 WP
Screen technology	TFT Color LCD
Active screen size (diagonal)	542 mm (21.5")
Active screen size (H x V)	267,3 mm x 475,2 mm (10.52" x 18.71")
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Resolution	2MP (1920 x 1080)
Pixel pitch	0.2475
Color imaging	Yes
Gray imaging	Yes
Color support	10 bit; 16,7 million colors
Viewing angle (H, V)	178°
Ambient Light Compensation (ALC)	Yes, preset values in OSD
Front sensor	Yes
Maximum luminance	250 cd/m² typical
DICOM calibrated luminance	180 cd/m²
Contrast ratio	1000:1 typical
Response time (Tr + Tf)	14 ms
Housing color	White + Silver

Video input signals	VGA, DisplayPort, HDMI
USB ports	1 upstream, 2 downstream
USB standard	2.0
Power requirements (nominal)	100-240 V
Power consumption (nominal)	40 W (<0,5 W in stand-by)
Power save mode	Yes
Power management	DPMS
Dot clock	165 MHz
OSD languages	English, French, German, Spanish, Italian, Dutch, Japanese, Traditional Chinese, Simplified Chinese, Korean
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Dimensions w/o stand (W x H x D)	507 x 306 x 57 mm
Dimensions packaged (W x H x D)	610 x 505 x 230 mm
Net weight with stand	6.7 kg
Net weight w/o stand	4.3 kg
Net weight packaged with stand	10.1 kg
Height adjustment range	80 mm
Tilt	0° / +20°
Swivel	-30° / +30°
Pivot	N/A
Mounting standard	VESA (75 mm)
Screen protection	Protective, non-reflective glass cover
Certifications	CE (MDD 93/42/EEC class I product), CE - 2004/ 108/EC, IEC 60950-1:2005 + A1:2009 (2ND EDITION), IEC 60601-1 2ND ED:1988 + A1:1991 + A2:1995, UL 60601-1 1ST EDITION, CAN/CSA-C22.2 NO. 601.1-M90:2005, IEC 60601-1:2005 + A1:2012, ANSI/AAMI ES 60601-1:2005 + C1:2009 + A1:2012, CAN/CSAC22.2 No. 60601-1-08:2008, DEMKO - EN 60601- 1:2006, EN 60601-1-2:2007, CCC - GB9254-2008 + GB4943-2001 + GB17625.1-2003, KC, VCCI, FCC class B, ICES-001 Level B, FDA Class I device, RoHS
Supplied accessories	User Guide
	Cable routing strap
	Cleaning cloth
	Video cables (1 x VGA + 1 x HDMI)
	Main cables (UK, European (CEBEC/KEMA) or USA (UL/ CSA; adaptor plug NEMA 5-15P) or Chinese (CCC))
	USB 2.0 cable
	External power supply (model: BridgePower Corp. BPM060S12F15; Input: 100-240 V AC, 50-60 Hz, 1.5 A; Output: +12V DC ($\overline{}$), 5.0 A)
Optional accessories	Proximity sensor
QA software	MediCal QAWeb 2
Warranty	3 years
Operating temperature	10°C - 40°C
Storage temperature	-20°C - 60°C
Operating humidity	30% - 75% (non-condensing)
Storage humidity	5% - 85% (non-condensing)

Operation altitude	2000 m
Storage altitude	7500 m
Operating pressure	70 kPa - 106 kPa
Storage pressure	50 kPa - 106 kPa