



Veraview X800

Operation Instructions

Panorama and CT

CAPC



Thank you for purchasing the Veraview X800.

For optimum safety and performance, read this manual thoroughly before using the unit and pay close attention to warnings and notes.

Keep this manual in a readily accessible place for quick and easy reference.

Registered Trademarks and Brand Names:

The names of companies, products, services etc. used in this manual are either registered trademarks or brand names.

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1 Prevent Accidents

1.1 Attention Customers

Do not fail to receive clear instructions concerning the various ways to use this equipment as described in this accompanying Operation Instructions.

Fill out and sign the warranty and give the dealer from whom you purchased the equipment a copy.

1.2 Attention Dealers

Do not fail to give clear instructions concerning the various ways to use this equipment as described in this accompanying Operation Instructions.

After instructing the customer in the operation of the equipment, have him fill out and sign the warranty. Then fill in your own section of the warranty and give the customer his copy. Do not fail to send the manufacturer's copy to J. MORITA MFG. CORP.

1.3 Prevent Accidents

Most operation and maintenance problems result from insufficient attention being paid to basic safety precautions and not being able to foresee the possibilities of accidents.

Problems and accidents are best avoided by foreseeing the possibility of danger and operating the unit in accordance with the manufacturer's recommendations.

First thoroughly read all precautions and instructions pertaining to safety and accident prevention; then, operate the equipment with the utmost caution to prevent either damaging the equipment itself or causing bodily injury.

The following symbols and expressions indicate the degree of danger and harm that could result from ignoring the instructions they accompany:

MWARNING

A hazard with a medium risk of injury or death if not avoided.

^CAUTION

A hazard with a low risk of minor or moderate injury if not avoided.

(Usage Note)

This informs the user of important points concerning operation or the risk of equipment damage.

The user (i.e., healthcare facility, clinic, hospital etc.) is responsible for the management, maintenance and use of medical device.

Disregarding the information on safety is considered ABNORMAL USE.

This equipment must only be used by dentists and other legally licensed professionals.

Do not use this equipment for anything other than its specified dental X-ray exposure purpose.

1.4 Disclaimer

- J. MORITA MFG. CORP. will not be responsible for accidents, equipment damage, or bodily injury resulting from:
 - 1. Repairs made by personnel not authorized by J. MORITA MFG. CORP.
- 2. Any changes, modifications, or alterations of its products.
- 3. The use of products or equipment made by other manufacturers, except for those procured by J. MORITA MFG. CORP.
- 4. Maintenance or repairs using parts or components other than those specified by J. MORITA MFG. CORP. and other than in their original condition.
- 5. Operating the equipment in ways other than the operating procedures described in this manual or resulting from the safety precautions and warnings in this manual not being observed.
- 6. Workplace conditions and environment or installation conditions which do not conform to those stated in this manual such as improper electrical power supply.
- 7. Fires, earthquakes, floods, lightning, natural disasters, or acts of God.

The useful life of the Veraview X800 is 10 years (based on self-certification) from the date of shipment provided it is regularly and properly inspected and maintained.

J. MORITA MFG. CORP. will supply replacement parts and be able to repair the product for a period of 10 years after the manufacture of the product has been discontinued. For the duration of this period, we will supply replacement parts and be able to repair the product.

1.5 Fill Out Safety Instruction Record

When the unit is installed or when there is a new user, the installer or other responsible party must explain the safety note in the Operation Instructions to the new user and the person responsible for maintenance and management, and fill out the Safety Instruction Record provided with the contents of the explanation, the names of the person responsible for maintenance and management, the new user, the person who made the explanation and other required items.

1.6 In Case of Accident

If an accident occurs, the Veraview X800 must not be used until repairs have been completed by a qualified and trained technician provided by the manufacturer.

1.7 User Qualifications

Intended Operator Profile

- a) Qualification: Legally qualified person such as radiologic technician and dentists for X-ray device operation (it may differs among countries).
- b) Education and Knowledge: It is assumed that the user understands the risks of X-rays and the protective steps required. It is also assumed that the user is thoroughly familiar with X-ray diagnosis, anatomy, and hygiene including the prevention of cross contamination.
- c) Language Understanding: English (Intended for professional use as described above)
- d) Experience: Experienced person with operating panoramic X-ray device.

No special training is required except in cases where this is required by legal regulations of the relevant country or region.

2 Precautions

MWARNING

- To avoid the risk of electric shock, this equipment must only be connected to the supply mains with protective earth.
- Do not modify this equipment without authorization of J. MORITA MFG. CORP.

Infection Control

- Proper infection control procedures must be established and maintained for each patient.
- Mouthpieces, disposable covers for the chinrest and bite-block must never be re-used. A new one must be used for each patient.
- The chinrest, bite-block, and upper lip rest must be disinfected by wiping with Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) after each use, and disposable covers put on the chinrest and bite-block.
- The patient handle must be disinfected by wiping it with Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) after each patient. If the patient has an open or bleeding wound on his hand, the patient handle should be covered to prevent cross contamination before the patient grips it.

MIMPORTANT PRECAUTIONS

These caution remarks are especially critical for safe operation and use.

- This unit must be used in a X-ray controlled area. Local regulation for radiation protection must be observed.
- Control box and emission switch must install and in radiation protected area.
- Special attention must be paid to the strength of the floor and wall materials, refer to the installation instructions.
- Steps must be taken to limit entry to the X-ray room during X-ray emission.
- User must restrict access to the equipment in accordance with local regulations for radiation protection.
- This equipment has an optional function so that it can be connected to external electrical devices such as a door switch that will stop X-ray emission or prevent X-ray emission. In this case we recommend that the management of the facility post a visible notice to inform people of the use of the external electrical devices.
- The patient must be provided with adequate X-ray protection.
- No unauthorized personnel be in the immediate area when the equipment is in use.
- If, for some reason, a person other than the patient must be within the X-ray controlled area during X-ray emission, he must take adequate X-ray protection measures.
- Proper radiation safety precautions must be established in accordance with local, state and governmental regulations in regards to operator and patient protection. The ultimate responsibility lies with the owner/operator to ensure that the protection requirements of national and local codes are met.
- Do not use this unit for fluoroscopic examinations.
- Implanted pacemakers and defibrillators may be adversely affected by continuous exposure to an X-ray beam. If an examination requires this, the operator must take proper care after referring to the critical safety remarks and interaction effects sections in the manual for the pacemaker or defibrillator.
- Do not use the wireless transmission devices listed below in the examination area:
 - (1) Cell phone terminals.
 - (2) Wireless transmitting devices such as ham radios, walkie-talkies, and transceivers.
 - (3) Personal Handy-phone System (PHS).
 - (4) Routers for intra-building paging systems, wireless LAN, cordless analogue telephones, and other electric wireless devices.

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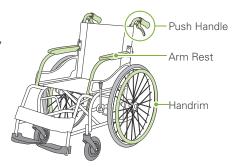
- Interference from the Veraview X800, devices listed below might malfunction or operate in a random, unexpected and dangerous manner.
 - (1) Electric medical devices for examination, diagnosis and treatment.
 - (2) Personal computers.
- Responsible organization in medical institution need for providing means for audio and visual communication between the operator and the patient.
- The operator must be able to see the exposure emissions lights and hear the audible signal during operation of the equipment.
- The operator must be able to see and hear the patient during the operation of the equipment.
- Watch the area around the moving parts to avoid collision against the body parts or other objects which may result in injury.
- Make sufficient space around the power distribution panel so that it is easily to breaker open in case of emergency.
- Do not open or remove any of the unit's enclosures. No user serviceable parts inside.
- The operator shall not touch the LAN connector when touching the patient.
- Do not use this equipment for patients when it is being maintained or serviced.
- CT images cannot be used for diagnosis of soft tissue.
- CT exposures are only appropriate when sufficient diagnostic information cannot be obtained with conventional intraoral radiographs.
- For optimum radiation protection, refer to the recommendations and advice issued by the ICRP (International Commission on Radiological Protection) and EADMFR (European Academy of Dentomaxillofacial Radiology).
 - * As Low As Reasonably Achievable (ALARA): All exposure to radiation should be kept at the lowest level as is reasonably possible taking into consideration social and economic factors. (ICRP, 1977)
 - * Basic Principles on the use of Cone Beam CT (EADMFR)
- This equipment does not provide CT Numbers (required in 21 CFR 1020.33 and IEC60601-2-44) or any equivalent values calculated from the absolute X-ray absorption of the target material.

Pregnant Women, Women in Labor, Nursing Mothers, and Infants

- Careful and prudent judgment must be exercised before exposing pregnant women, women who might be pregnant, women in labor, nursing mothers, infants etc. to X-ray examination, the X-ray dosage should be reduced as much as possible and the examinee must be provided with adequate X-ray protection.
- Judgment and caution should be used in regards to radiographs of pregnant women. The decision should be based on "clinical need of diagnostic information".

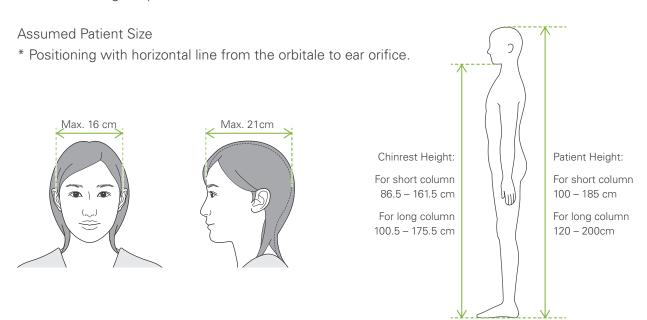
Patient in a Wheelchair or Sitting in a Chair

- This unit has an electric lift mechanism. Whenever the lift is lowered, make sure none of its parts (control panel support, arm, arm support, cephalo unit, cephalo support) hit or contact the patient.
- Make sure the push handles, arm rests, handrims etc. of a wheelchair do not make contact with the X-ray unit.
- Before making an exposure, run a test with no X-ray mode to make sure that the patient or the wheelchair does not make contact with the arm.

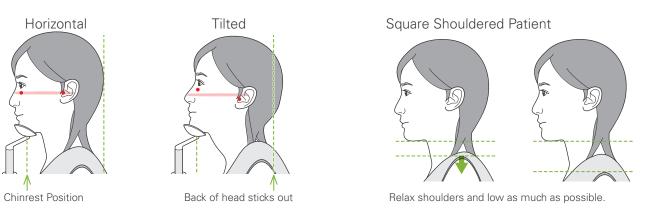


Patient Size and Positioning

- The X-ray detector, X-ray head and arm rotate around the patient during emission. Depending on the patient's head size, positioning and shoulder location, the arm might make contact with the patient. To avoid this, the operator must carefully consider the patient's size and positioning.
- If the patient might be larger than permitted, run a test with no X-ray mode to make sure that the arm does not make contact with the patient.
- For a CT exposure of a large or larger than permitted patient, use 180° exposure mode to keep the arm from contacting the patient.



- If the line from the orbitale to the ear orifice is not horizontal, the back of the head might stick out and make contact with the arm. If this is the case, run a test with no X-ray mode to make sure the arm does not contact the patient.
- For patients who have square shoulders or short necks, the arm might contact the shoulder. In this case, have the patient lower his shoulders as much as possible and then run a test with no X-ray mode to make sure the arm does not contact the patient.



3 Production Information

3.1 Capabilities

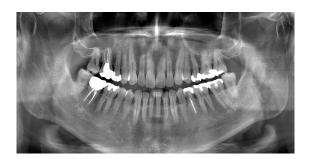
The Veraview X800 is a digital dental X-ray equipment that can make panoramic image-layer exposures, cone-beam CT exposures (hereinafter referred to as CT), and cephalometric exposures (option).

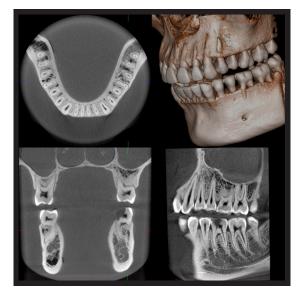
Panoramic Exposures

- Dental Arch Panorama (Standard, Shadow less, Ortho)
- Maxillary Sinus Panorama (Anterior, Posterior)
- Quadruple TMJ (Standard, Linear)
- Bitewing

CT Exposures

- Dental Arch
- TMJ
- Maxillofacial





Cephalometric Exposures (option)

- Lateral
- PA (posterior-anterior)
- 45° Angle
- Hand



3.2 Related Documents

To use this unit, read the documents listed below and follow all cautionary notes and usage methods.

• Operation Instructions

3.3 Acronyms

FOV: Field of View, mainly for CT exposure regions.

ROI: Region of Interest (exposure areas)

3.4 Environmental Considerations

Disposal of this unit, its accessories and replacement parts must conform to all local and international regulations.

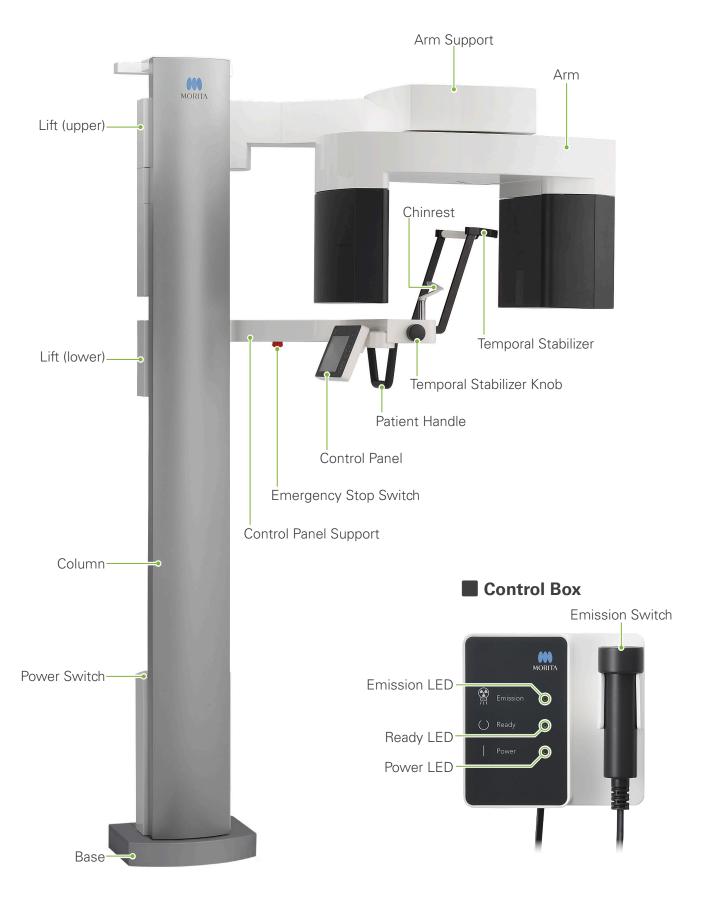
Materials used in this unit:

• X-ray Head: Lead, High pressure dielectric oil

X-ray Detector: LeadCollimator: Lead

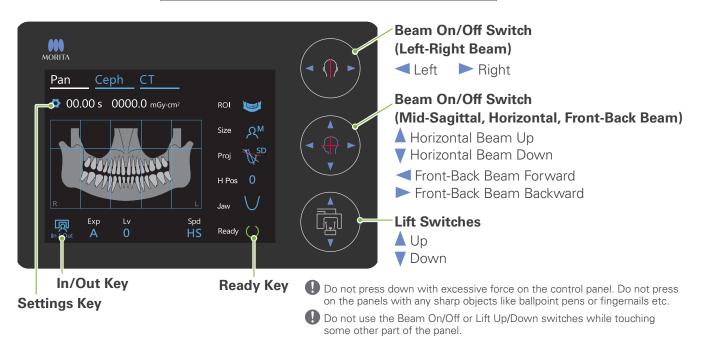
4 Parts Identification

■ Main Unit





Control Panel Blue icons and the Ready key are activated by touch.



• Mouthpieces (1 box of 50 pcs.) • Chinrest • Chin Rest (1) • Bite-Block (1) • Upper Lip Rest (1) • Chinrest Disposable Cover (1 box of 100 pcs.) • Bite-Block Cover (1 box of 300 pcs.)

5 Before and After Use

5.1 Operation Conditions

Use Veraview X800 Under Following Conditions:

Temperature: +10°C to +35°C (+50°F to +95°F); Humidity: 30% to 75% (without condensation); Atmospheric Pressure: 70 kPa to 106 kPa

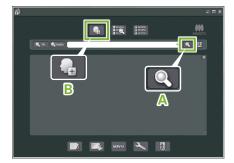
- * If an accident occurs, the equipment must not be used until repairs have been completed by a qualified and trained technician authorized by the manufacturer.
- * Have patients remove glasses, earrings and any other accessories that might interfere with the exposure.
- * Before use make a start-up inspection to make sure the unit will operate properly and safely.

5.2 Set Up

5.2.1 Start i-Dixel

1. Start i-Dixel

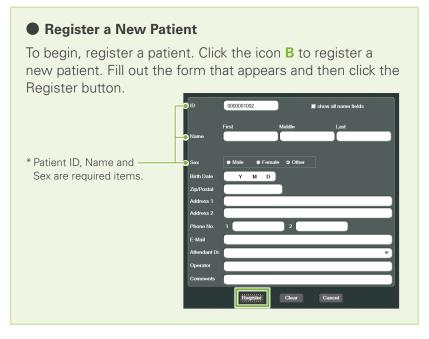
Start the i-Dixel software.





2. Retrieve a registered patient

Enter the patient ID in the search field and click the search icon (A). To use the patient list, select a patient and then double-click it.

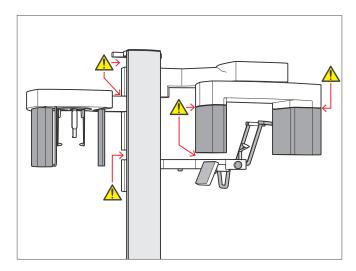


An image list will appear.

* In CT exposure mode, the resolution is set automatically and cannot be changed.



5.2.2 Safety Check for Moving Parts

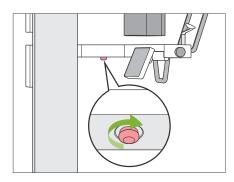


For safety, make sure that fingers will not be pinched before moving any parts.

ACAUTION

• Keep fingers away from gaps and openings of moving parts as well as the holes on the support column.

Emergency Stop Switch



In Case of Emergency

Press the Emergency Stop switch. This will stop the arm rotation and X-ray emission.

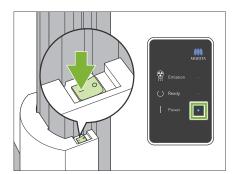
* Never use this switch except in the case of an emergency.

After Pressing Emergency Stop switch.

- 1. Guide the patient away from the unit and turn off the Power switch.
- 2. Restore the unit to a safe condition.
- 3. Turn the Emergency Stop switch in the direction shown by the arrow in the illustration until it clicks to release it.
- 4. Restart the computer.
- 5. Turn on the Power switch.
- 6. Check the panoramic, CT, and cephalometric exposure operation.

If the unit cannot be returned to a safe condition or will not operate, contact your local dealer or J. MORITA OFFICE.

5.2.3 Start Main Unit



Press the side of the Power switch on the lower part of the column that is marked with a line (|).

The unit will turn on and the blue Power LED on the control box will light up.

ACAUTION

- Do not press the Power switch if the patient is near or inside the equipment. Otherwise, the arm will rotate and the control panel support will move and could injure the patient.
- If the Emergency Stop switch has been inadvertently pressed while the unit was being cleaned etc., the unit cannot be turned on. Release the Emergency Stop switch, and then turn on the unit.
- (1) When starting up the Veraview X800 right after turning it off (e.g., rebooting the equipment), wait at least 5 seconds to turn it on again.

The "PC and cassette communication check" dialogue box will appear.



CAUTION

• If the temporal stabilizers are open, a message will appear warning to check them. The arm might hit them so turn the knob and close them.

PC and cassette communication check

Please wait while communication between the digital cassette and PC is verified.

Initialization

The unit will move to its initial position. The arm will move in multiple directions so ensure the immediate area is clear of obstructions before proceeding. If a patient is positioned in the unit, have them exit the unit before proceeding.



Then the "Initialization" dialogue box will appear.

If the patient is already in position, have him step away from the equipment.

Check the area and then click the "Ok" button.

The arm will move to the patient entry position.

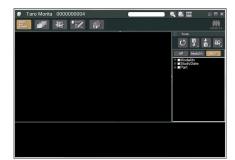


• Do not fail to move the patient away from the equipment and check the area for safety before initializing the equipment. Otherwise, the arm might move and it could hit the patient.

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5.3 Start-up Inspection

Before use make a start-up inspection to make sure the unit will operate properly and safely.



1. Open the image list for the start-up inspection

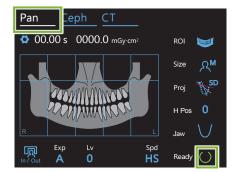
Select the test patient used for the start-up inspection from the i-Dixel patient list and display the image list for it.

● First Time Only

To begin, the test patient must be registered in i-Dixel.

Click the Patient Registration Window icon to display the registration window, fill out the form and then click the Register button.







2. Check operation for panoramic exposure

Touch the Pan key on the control panel to set the unit for panoramic exposure mode.

Set the exposure conditions as shown below.

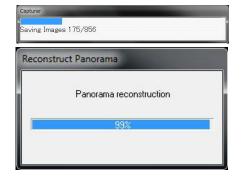
- Exposure Setting (Exp): "M" (manual)
- Tube Voltage (kV): 60 kV
- Tube Current (mA): 2 mA

Touch the Ready key to light it up.

Hold down the Emission switch and check the following items:

- Arm moves and X-rays are being emitted.
- The Emission LED is on and the melody is playing.
- X-rays emission and the arm rotation stop when the exposure is completed.
- The Emission LED turns off and the melody stops.

Release the Emission switch and hang it in its holder on the control box.



3. Panoramic image display

The "Reconstruct Panorama" window will appear in i-Dixel, and after about 10 seconds the image will appear.



A two-toned beep sound when transmission has been completed and the LED turns green and blinks, but another exposure cannot be made until the image appears in the computer screen.



4. Check operation for CT exposure

Touch the CT key on the control panel to set the unit for CT exposure mode.

Set the exposure conditions as shown below.

- Exposure Setting (Exp): "M" (manual)
- Tube Voltage (kV): 70 kV
- Tube Current (mA): 2 mA

Check the unit's operation in the same way as for a panoramic exposure.



5. CT image display

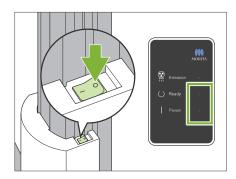
The inspection image will appear after about 5 minutes.

6. Delete the inspection data

Delete the data acquired during the inspection exposure.

5.4 After Use

5.4.1 Turn Off the Unit



Press the side of the Power switch marked with a circle (). The unit will turn off. The Ready and Power LEDs will turn off.

CAUTION

• Do not fail to turn off the Power switch. This prevents the risk of current leakage, inadvertent operation, etc.

When starting up the Veraview X800 right after turning it off (e.g., rebooting the equipment), wait at least 5 seconds to turn it on again.

6 Panoramic Exposures

6.1 Exposure Types and Functions

6.1.1 Exposure Areas and Projections

Dental Arch

This is a single image of the whole dental arch. There are three possible projections.



The most common type.







Shadow Less

Reduces shadows obscuring the mandibular ramus.







Ortho

This aims the X-ray beam at right angles to the dental arch to reduce the overlapping of individual tooth images.





AFP: Adaptive Focal Point

This function finds the optimum focus for every position from the data about layers obtained by the image-layer exposure. Everything in the image from the root apex to the incisor region is in focus.

AGS: Adaptive Gray Scale

This automatically adjusts density to make the whole panoramic image clearing observable including the dental arch, jaw bone, TMJ etc.

AIE-HD: Auto Image Enhancement - High Definition

Optimizing panoramic image processing makes every detail sharp and clear.

* The AFP, AGS, and AIE-HD functions are operated on i-Dixel. For details refer to the Operation Instructions for i-Dixel.

■ Maxillary Sinus

A panoramic image mainly used to examine the maxillary sinus and facial injuries. There are two types, anterior and posterior.



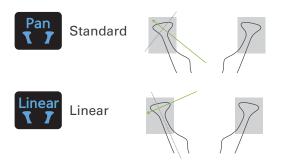


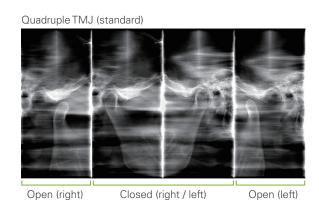




Quadruple TMJ

Four digital images will appear in the computer display: one each for the mouth open and closed on both sides. The X-ray beam angle is optimum for the average distance between the joints and the average length of the image layer.





Bitewing

For occlusal exposure. Useful for diagnosis of mild periodontitis or caries in the proximal spaces of premolars and molars and for prosthetics as well.











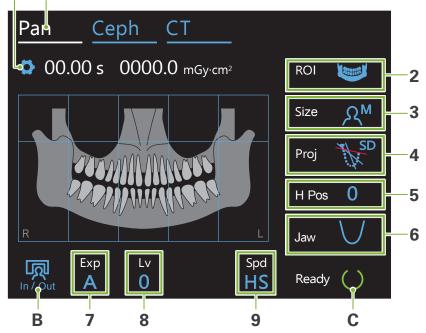
6.1.2. DDAE (Digital Direct Auto Exposure)

During the exposure the flat panel detector detects X-ray transparency in real time and then controls the amount of X-rays emitted to create images with much better dynamic range. Contrast can be adjusted with the auto level value.

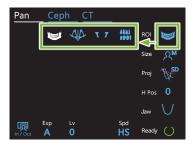
6.2 Operation and General Settings

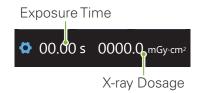
6.2.1 Panoramic Exposure

A 1. Panoramic Exposure Mode Key



Numbers 2 to 9 show the current settings. Touch any one of the icons to show other selections.





A. Settings Key

Hold down this key to save the current settings.

B. In/Out Key

Touch for patient entry and egress. The arm moves to 90° to the lift for easy patient entry and egress.

C. Ready Key

Shows that the unit is in the ready state.

This key blinks when the Power switch is turned on.

Touch it after guiding the patient into position: it will light up. The arm will move to its patient position and the mid-sagittal, horizontal, and front-back beams will light up. The unit is ready to begin X-ray exposure whenever the Emission switch is pressed.

6.2.2 Positioning Beams

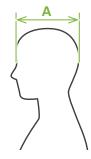


6.2.3 Settings

2.	3.	4.	5.	6.	7.		8		0
Region	Patient Size	Projection	Arm Height	Dental Arch Shape	Exposure Setting	If No. 7 is auto	If No. 7 is	s manual VI	9. Scan Speed
(ROI)	(Size)	(Proj)	(H Pos)	(Jaw)	(Exp)	(Lv)	(kV)	(mA)	(Spd)
		Standard	15	Standard					
Dental Arch	(AC	Shadow Less	15	Narrow		+6			
		SL Ortho	10	Wide	A (auto exposure)	+5		10*	
		OT	5	VVIde		+4	90	9	
	ر کاری	Anterior	0		M	+3	85	8	HD
Maxillary Sinus	عد	The state of the s	U		(manual exposure)	+2	80	7	(standard speed, high definition)
		Posterior	mm		Off	+1	75	6	
	$\mathcal{L}_{\mathbf{M}}$	Standard			(no X-ray emission)	0	70	5	HS
TMJ		Pan T				-1	65	4	(high speed)
		Linear Linear				-2	60	3	
	\mathcal{L}_{Γ}		/	/	M	-3		2	
Bitewing		Bitewing BW	15 10 5 0		(manual exposure) Off (no X-ray emission)	-4			

^{*} For 85 or 90 kV, mA setting is from 2 to 9.

■ Patient Size



The set values for patient size is the sagittal length of the skull (A).

Also the upper limit for the lift cannot accommodate patients taller than 180 cm. We recommend having patients who are too tall to sit down.

		C (Child)	S (Small)	M (Middle)	L (Large)
Sagittal Leng	th	max. 17 cm	max. 17 cm	max. 19 cm	19 – 21 cm

^{*} The above values are estimates only. The dentist should judge this by the patient's body shape, skeletal structure etc.

Child Mode

The arm rotation angle is smaller to make the irradiation range narrower and reduce X-ray dosage.

* Child mode is for children and patients who have a small jaw bone. If the jaw bone is too large, the TMJ may not appear in the image. For the entire jaw to appear in the image, the length of a horizontal line from the center of the incisors to the outer ear orifice should be less than 7 cm.



Child Mode Panoramic Image

Emission Settings

Setting the level for auto exposures "A" at "0" will reduce the X-ray dosage. Adjust the auto level depending on the desired result such as increasing contrast or getting a smoother image.

Reference Settings

A (auto exposure): +2 or above

M (manual exposure): 75 kV, 8 mA or above





Auto Level "+2" Image

Auto Level "0" Image

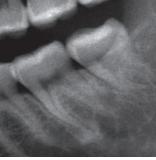
Scan Speed

"HS" (high speed) takes about half the time as "HD" (standard speed, high quality). Refer to the information below, and select the one that suits your purposes.

HS: Use this for patients who find it hard to stand still or to reduce X-ray dosage.

HD: This produces images with better contrast and less noise. X-ray dosage is greater than "HS".



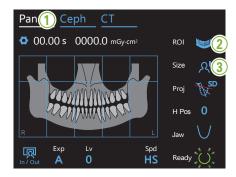


HD Image

HS Image

6.3 Panoramic Exposure Methods (Dental Arch, Maxillary Sinus, and Bitewing)

6.3.1.1 Preparation (Dental Arch and Maxillary Sinus Exposures)

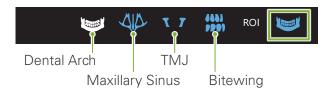


1. Select panoramic exposure mode

Touch the Pan key to set the unit for panoramic exposure mode.

2. Select region of interest (ROI)

Touch the icon to the right of "ROI" to select the region of interest.



3. Select patient size

Touch the icon to the right of "Size" to select the patient size.

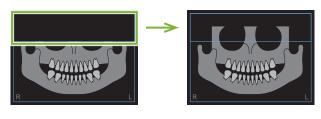


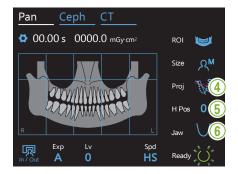
Child Mode

The factory setting for the upper part of the image is off.

Touch the square at the top of the illustration to add this part to the exposure area.

* Only for dental arch panorama.





4. Select projection

Touch the icon to the right of "Proj" to select a projection.

Dental Arch Panorama

SD (standard): Standard panorama

SL (shadow less): Reduces shadows obscuring the mandibular ramus.

OT (ortho): Reduces overlapping tooth images.



Maxillary Sinus Panorama



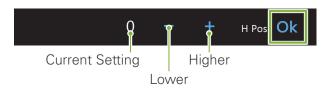
5. Adjust arm height

* If necessary.

This is in case the arm makes contact with the patient's shoulder.

- If the arm is raised, the location of the exposure will be higher and the end of the chin may not be exposed. First have the patient lower his shoulders as much as possible. If the arm still makes contact with the shoulders, adjust the arm height with H Pos.
- If the lift reaches its upper limit, a series of beeps will sound, and the arm cannot be raised any more.

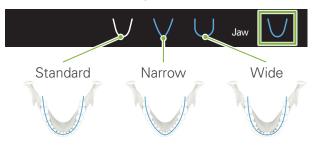
Touch the icon to the right of "H Pos" to adjust the arm height. It can be set from 0 to 15 mm in 5 mm increments.

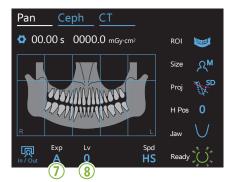


6. Select dental arch shape

* Only for dental arch panorama.

Touch the icon to the right of "Jaw" to select the dental arch shape.





7. Set exposure mode

Off

Α

Touch the letter under "Exp" to set the exposure mode.

Auto Exposure (digital direct auto exposure function)
Depending on the patient type and exposure area,
X-ray emission is monitored and adjusted in real time.
Contrast can be adjusted using the auto level value.

Manual Exposure

Set tube voltage (kV) and current (mA) manually.

No X-ray Emission

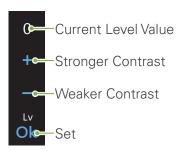
This function is prepared to simulate an actual scan without X-ray emission.

Use it to check the arm rotation orbit, to explain the movement of the device to the patient or to ensure the arm will not hit the patient during a scan.

* If "A" (auto exposure) was selected in step 7.

8-A. Set auto level

Touch the number under "Lv" to set the auto level value.



X-ray dosage will change depending on the auto level value.

If X-ray dosage is increase, image contrast and density will increase and hard tissues will be clearer and sharper. If the X-ray dosage is decreased the image quality for soft tissue will be better.

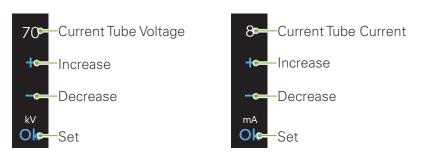
Auto Level and Air Kerma

Auto Level	Air Kerma Ratio (X-ray Output Ratio)	Image Quality Applicability
+6	1.40	Hard Tissue
+5	1.33	hard fissue
+4	1.26	
+3	1.19	
+2	1.12	
+1	1.06	
0	1	
-1	0.94	
-2	0.88	
-3	0.83	♦
-4	0.77	Soft Tissue

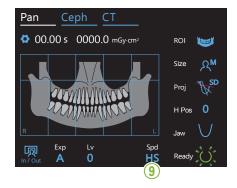
* If "M" (manual exposure) was selected in step 7.

8-M. Set tube voltage (kV) and current (mA)

Touch the numbers under "kV" and "mA" to set the values. Tube voltage can be set from 60 - 90 kV in 5 kV increments. Tube current can be set from 2 - 10 mA in 1 mA increments.



Patient Size	С	S	М	L
Estimated Tube Voltage (kV)	75	75	75	75
Estimated Tube Current (mA)	6	8	8	8



9. Select scan speed

Touch the letters under "Spd" to set the scan speed.

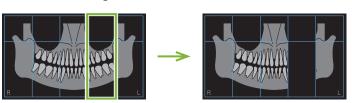
HD—Standard Speed, High Definition

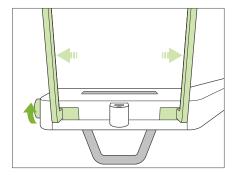
HS—High Speed (reduces exposure time by about half)

10. Select non-exposure area

* For dental arch panorama only. If necessary.

To reduce the X-ray dosage, a partial exposure can be made. Touch the rectangular areas in the panoramic image to make them black and stop them from being exposed to X-ray emission. Press the area again to restore it.





11. Set chinrest or bite-block

Turn the temporal stabilizer knob and open the stabilizers.

Wipe the chinrest or bite-block with Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) and put it in the chinrest holder.

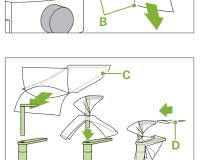
* Before use, make sure the component is not scratched or damaged in any way.

Chinrest

- 1. Open a disposable cover and put it on the chinrest.
- 2. Pull down the paper and peel it off
- A: Cover (semi-transparent)
- B: Paper (white)

Bite-Block

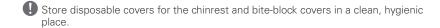
- Peel off the paper of a chinrest disposable cover and cover the bit (the part of the bite-block the patient bites on) with it.
- Twist the disposable cover and put the bit in the bite-block stand.
- 3. Put a bite-block cover on the hit



- C: Chinrest Disposable Cover
- D: Bite-block Cover

MWARNING

- The chinrest and bite-block must be disinfected by wiping with Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) after each use, and before putting on a disposable cover.
- Disposable covers for the chinrest and bite-block must never be re-used. A new one must be used for each patient.
- The patient handle must be disinfected by wiping it with Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) after each patient. If the patient has an open or bleeding wound on his hand, the patient handle should be covered to prevent cross contamination before the patient grips it.

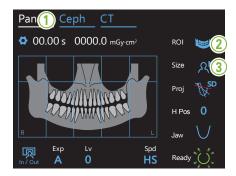




To continue go to "6.3.2 Patient Entry and Positioning" (p. 34).

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6.3.1.2 Preparation (Bitewing)



1. Select panoramic exposure mode

Touch the Pan key to set the unit for panoramic exposure mode.

2. Select region of interest (ROI)

Touch the icon to the right of "ROI" to select the region of interest.



3. Select patient size

Touch the icon to the right of "Size" to select the patient size.





4. Select areas to trim

* For dental arch panorama only. If necessary.

To reduce the X-ray dosage, a partial exposure (left or right area only) can be made. Touch one of the square areas as shown in the illustration below to make it black and stop it from being exposed to X-ray emission. Press the area again to restore it.



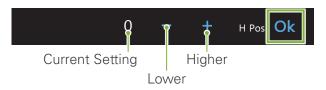
5. Adjust arm height

* If necessary.

This is in case the arm makes contact with the patient's shoulder.

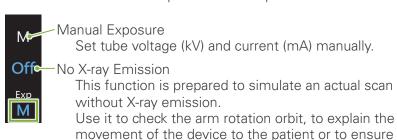
- If the arm is raised, the location of the exposure will be higher and the end of the chin may not be exposed. First have the patient lower his shoulders as much as possible. If the arm still makes contact with the shoulders, adjust the arm height with H Pos.
- If the lift reaches its upper limit, a series of beeps will sound, and the arm cannot be raised any more.

Touch the icon to the right of "H Pos" to adjust the arm height. It can be set from 0 to 15 mm in 5 mm increments.



6. Set exposure mode

Touch the letter under "Exp" to set the exposure mode.

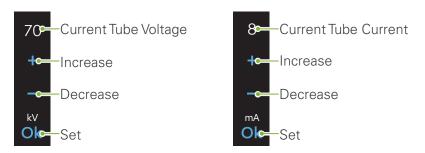


the arm will not hit the patient during a scan.



7. Set tube voltage (kV) and current (mA)

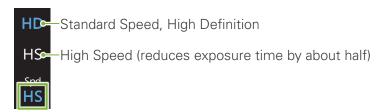
Touch the numbers under "kV" and "mA" to set the values. Tube voltage can be set from 60 - 90 kV in 5 kV increments. Tube current can be set from 2 - 10 mA in 1 mA increments.

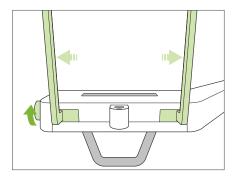


Patient Size	С	S	М	L
Estimated Tube Voltage (kV)	75	75	75	75
Estimated Tube Current (mA)	6	8	8	8

8. Select scan speed

Touch the letters under "Spd" to set the scan speed.





9. Set chinrest or bite-block

Turn the temporal stabilizer knob and open the stabilizers.

Wipe the chinrest or bite-block with Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) and put it in the chinrest holder.

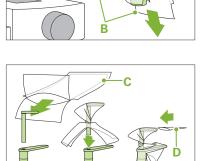
* Before use, make sure the component is not scratched or damaged in any way.

Chinrest

- 1. Open a disposable cover and put it on the chinrest.
- 2. Pull down the paper and peel it off.
- A: Cover (semi-transparent)
- B: Paper (white)

Bite-Block

- Peel off the paper of a chinrest disposable cover and cover the bit (the part of the bite-block the patient bites on) with it.
- Twist the disposable cover and put the bit in the bite-block stand.
- 3. Put a bite-block cover on the bit



C: Chinrest Disposable Cover

D: Bite-Block Cover

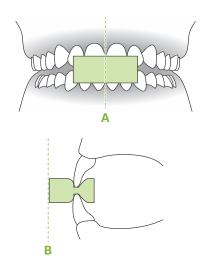
MWARNING

- The chinrest and bite-block must be disinfected by wiping with Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) after each use, and before putting on a disposable cover.
- Disposable covers for the chinrest and bite-block must never be re-used. A new one must be used for each patient.
- The patient handle must be disinfected by wiping it with Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) after each patient. If the patient has an open or bleeding wound on his hand, the patient handle should be covered to prevent cross contamination before the patient grips it.
- Store disposable covers for the chinrest and bite-block in a clean, hygienic place.

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6.3.2 Patient Entry and Positioning

For patient positioning, refer to the 2 Precautions, "Patient Size and Positioning" (p. 9).



1. Prepare patient

Have the patient wear an X-ray protection apron.

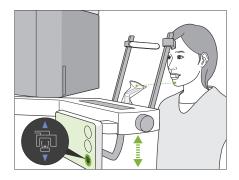
Have the patient bite on a new mouthpiece for incisal occlusion. Center the upper and lower incisors (A), and make the front surface of the mouthpiece perpendicular (B).

▲WARNING

• A new, uncontaminated mouthpiece must be used for each patient to prevent cross contamination.

ACAUTION

- The patient must wear an X-ray protection apron.
- Patient must remove glasses and all accessories like necklaces etc. Otherwise the exposure could fail.
- Use only the specified mouthpieces.
- Make sure the patient's hair cannot get caught in moving parts. If the patient's hair is tied in the back like a pony tail etc., have the patient undo it. Otherwise, the arm might hit the patient.
- Store mouthpieces in a clean, hygienic place.



2. Adjust unit height

Adjust the height of the unit so that the chinrest is even with the patient's chin.

Press and hold the Lift Up/Down switches to raise or lower the unit; release the switch to stop the unit.

ACAUTION

- Take care that the temporal stabilizers do not strike the patient in the eye.
- Take care that the patient is not hit or pinched by any part of the lift assembly (control panes support, arm, and arm support).
- Always use the Lift Up/Down switches to adjust the unit's height. Never try to force it; this would damage the unit.
- If the lift reaches its upper limit, a series of beeps will sound. If at this point the Ready key is touched, a message will appear saying that this is the limit for a CT exposure. If you make a CT at this point, the maxillary region may not be fully exposed. In this case, have the patient sit down or lower his position in some other way.

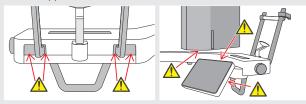


3. Patient entry

Have the patient step forward and put his chin on the chinrest. Have the patient lightly hook his thumbs on the patient handle.

↑ CAUTION

- Do not exert force to have the patient enter or leave; this could damage the temporal stabilizers.
- Never let the patient touch any of the control panel keys and switches.
- Keep fingers away from gaps and openings of moving parts as well as the holes on the support column.



• The chinrest can withstand a load of 2 kg and the patient handle and control panel support a load of 5 kg. These parts could be broken or the patient injured if the patient rests too much body weight on them.

Patient stands straight and neck is as perpendicular as possible.

If the patient stands (or sits) too far back, the upper torso will lean forward and the neck will be slanted. Have the patient move forward in this case.

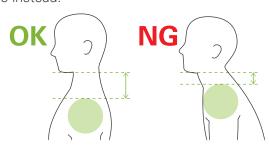
Pull jaw in and make the line from the orbitale to the ear orifice horizontal.

To get correct posture, it helps to tell the patient to move his forehead forwards while keeping his neck stretched while the lift is being lowered.

Relax and lower shoulders.

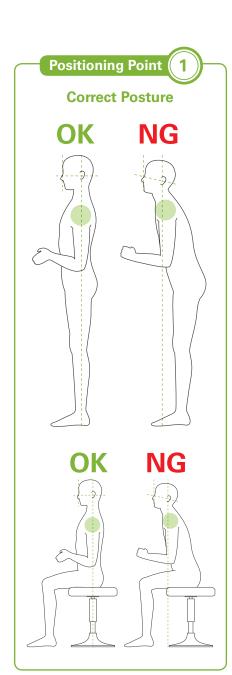
If the shoulders are too close to the jaw, the arm might hit them. For patients with square shoulders or a short neck, have them relax and lower their shoulders as much as possible.

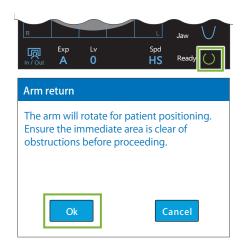
If the neck is slanted, have the patient straighten it and pull in his jaw. If the shoulders are too tension, the patient may be gripping the handle too tightly. Have patients lightly hook their thumbs on the handle instead.



▲CAUTION

• For patients who have square shoulders or short necks, the arm might contact the shoulder. In this case, have the patient lower his shoulders as much as possible and then run a test with no X-ray mode to make sure the arm does not contact the patient.





4. Touch Ready key

Touch the Ready key.

The "Arm return" dialogue will appear. Make sure the area is safe, and then touch the "Ok" button.

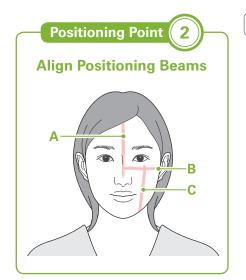
()

If a dialogue box asking to check the temporal stabilizers appears, touch the "Ok" button.

The arm will move to its patient position and the mid-sagittal, horizontal and front-back beams will light up.

MWARNING

•The laser beam could damage a person's eyesight; never look directly into the beam and let it strike anyone's eyes.





Horizontal Beam

▲ Up

Down

Front-Back Beam

Forward



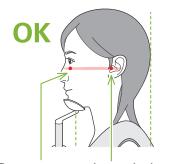
5. Line up beams

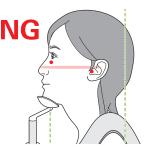
A: Mid-sagittal beam lines up with mid-sagittal plane.

Move the patient's head to line it up with the beam.

B: Horizontal beam lines up with the line from orbitale to ear orifice.

Have the patient keep a posture with the line from orbitale to ear orifice horizontal and then touch the Beam Up/Down switches to line the beam up with it.





Beam passes through these two points.

ACAUTION

• If the line from the orbitale to the ear orifice is not horizontal, the back of the head might stick out and make contact with the arm during an exposure or ready move. For a patient who has a large head, run a test with no X-ray emission mode to make sure the arm does not contact the patient.

C: Front-back beam lines up with the distal side of the left canine teeth.

First line the patient up so that the beam is roughly in the right position. Then press the Beam Forward/Backward switches to move the beam so that it is properly lined up.

- * If the beam cannot be line up with the distal side of the left canine teeth for edentulous patients or some other reason, line it up about 1 cm in back of the corner to the mouth
- * The front-back beam can be moved from -20 mm to +15 mm (less in some exposure modes). However, to make a panoramic scout exposure, moving the beam too much could make positioning less accurate.



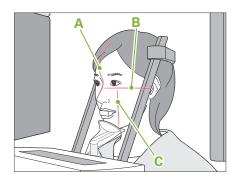
6. Close temporal stabilizers

Adjust the height of the temporal stabilizers.

Turn the temporal stabilizer knob and close the stabilizers firmly.

ACAUTION

• Do not use excessive force to close the temporal stabilizers. This could be uncomfortable for the patient or damage the stabilizers.



7. Check beams

Check the positioning of the all beams.

Mid-sagittal beam (A) lines up with mid-sagittal plane.

Horizontal beam (B) lines up with the line from orbitale to ear orifice. Front-back beam (C) lines up with the distal side of the left canine teeth.

Tell the patient not to move his head during X-ray emission (or while the melody is playing) and then leave the X-ray room.

ACAUTION

- Warn the patient not to move during emission (while the melody is sounding). If the patient moves, he could be hit by the arm or the image might be ruined.
- If the patient watches the arm go around, he may move his head. It is a good idea to tell the patient to close his eyes.

Min. Scan Move Function

(If this function is turned on.)

Before leaving the X-ray room, the arm can be moved to the position which will require the shortest exposure time (least time needed to hold down the Emission switch).

8. Touch Ready key

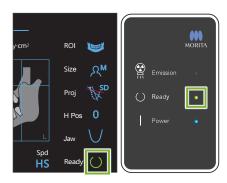
Make sure the patient has his hands on the patient handle. With the unit already in the ready state, press the Ready key again. The arm will move to its Min. Scan Move position and then a two-toned beep will sound.

Min. Scan Move:

- X-ray emission will begin as soon as the Emission switch is pressed.
- Only the Ready key will work.
- * Touch the Ready key again the arm will move back to its original patient positioning position.
- If after the arm moves to the Min. Scan Move position, one of the beams is not properly line up, touch the Ready key again to move the arm back to its original patient positioning position and re-position the patient.

After the arm moves, re-check the patient's positioning and hand position on the patient handle and then leave the X-ray booth.

6.3.3 Exposure



1. Check ready state

Check that the Ready key on the control panel and the Ready LED (green) on the control box are on.

2. X-ray emission



The arm will rotate and X-ray emission will start.

The Emission LED (yellow) on the control box will light up and the melody will sound.

* If both sides are selected to make a bite-wing exposure, two exposures will be made. Do not release the Emission switch before both are made.

MWARNING

• If an emergency occurs, release the Emission switch. Or press the Emergency Stop switch. This will stop the arm rotation and X-ray emission.

ACAUTION

- Go outside the X-ray room to press the Emission switch.
- If the operator must stay in the X-ray room for some reason, he should wear an X-ray protection apron and stay at least 2 meters away from the X-ray emission focal point. Also he should stay outside the area exposed to X-ray emission.
- Releasing the Emission switch during X-ray emission will stop the arm's rotation and terminate the exposure. In this case guide the patient away from the unit. If the X-ray head or the detector is directly in back of the patient, turn the unit off and slowly rotate the arm so the patient can leave. If you perform the ready procedures without having the patient move away from the unit, the arm could hit and possibly injure the patient.
- If an error occurs during an exposure, the arm will stop and the exposure will be terminated. In this case guide the patient away from the unit. If the X-ray head or the detector is directly in back of the patient, turn off the unit and slowly rotate the arm so the patient can leave. If you perform the ready procedures without having the patient move away from the unit, the arm could hit and possibly injure the patient.
- ① As much as 15 seconds could elapse from the time the Emission Switch is pressed until X-ray emission actually starts. This is not abnormal. It takes some time for the unit to verify the computer settings.
- If the computer is not ready, an error message will appear in the control panel. Check what the error is and turn off the X-ray unit. When the computer is ready, turn on the unit again.



3. Exposure completed

The melody stops when the exposure has been completed.

The arm will automatically move to the patient egress position.

Release the Emission switch and hang it in its holder on the control box.

6.3.4. Patient Egress



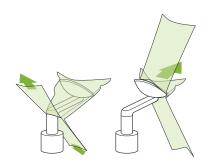
1. Guide patient away from unit

Turn the temporal stabilizer knob and open the stabilizers completely.

Guide the patient away from the unit.

ACAUTION

- Take care that the temporal stabilizers do not strike the patient in the eye.
- Be careful of the stabilizers when the patient leaves the unit.

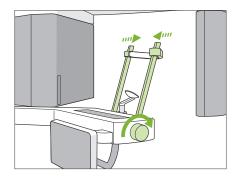


2. Throw away disposable covers etc.

Take the mouthpiece from the patient and throw it away.

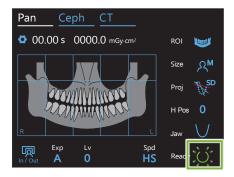
Throw away the chinrest or bite-block cover.

If the chinrest cover is removed by splitting its seams and turning it inside out, the outer surface will not touch anything.



3. Close temporal stabilizer

Turn the temporal stabilizer knob and close the stabilizers completely.



4. Touch Ready key

Touch the Ready key.

The "Arm return" dialogue will appear. Make sure the area is safe, and then touch the "Ok" button.

The arm will move back to its original patient positioning position.

* If another exposure will be made right away, touch the In/Out key to move the arm to its patient entry position.

6.3.5 Image Transmission



1. Image transmission

After the exposure is completed, the image is sent to i-Dixel. During transmission, the Ready LED is orange and blinks on and off.

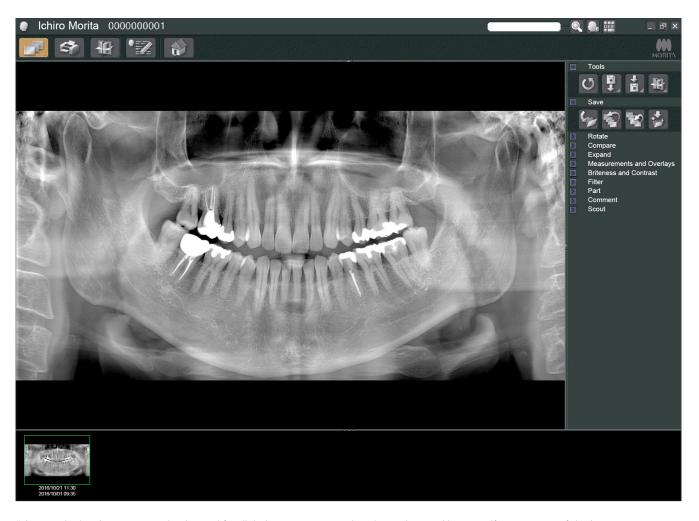


2. Image display

During image transmission, the "Capturer" window will appear in i-Dixel. Then the "Reconstruct Panorama" window with a progress bar will appear while the panoramic image is being reconstructed. The image will appear after about 10 seconds.

After transmission has been completed, the Ready LED on the control box and the Ready key in the control panel will turn green and blink on and off.

① A two-toned beep sound when transmission has been completed and the LED turns green and blinks, but another exposure cannot be made until the image appears in the computer screen.

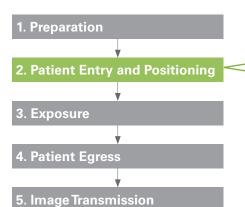


- * Automatic density compensation is used for digital exposures to make a better image. However, if some areas of the image are exceptionally dark, the density compensation will tend to make the entire image whiter than usual.
- * Depending on the X-ray exposure conditions and the patient's physiognomy, there may be a sudden, horizontal shift in density or light horizontal lines. This is not a malfunction or failure; it is due to subtle differences in sensitivity for sections of the Flat Panel Detector.
- * In areas with high X-ray opacity such as around Implants and prosthetics, a horizontal black line may appear. This is not a malfunction or failure; it is due to subtle differences in the individual photo diodes (pixels) on the Flat Panel Detector.
- * The depth of the image layer for a panoramic image can be slightly adjusted with i-Dixel (adjustable image layer panorama). If positioning was slight off, the focus can be corrected without having to retake the exposure. Focus correction can be done for the whole image or for upper and lower jaws separately.

6.3.6 Patient Positioning and Exposure Examples

Panoramic Exposures Flow Chart

■ Panoramic Exposure Points

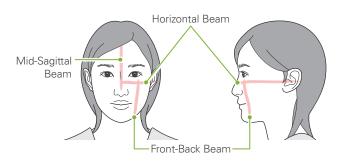


Important Step!!!

Patient positioning is the key to getting good panoramic images. Take special note of the "**Positioning Points**" in the foregoing instructions to make good exposures.

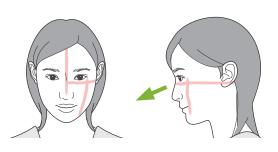
If patient positioning is not performed properly and accurately, the resulting image may be hard to use for diagnosis. Refer to the examples below to perform accurate patient positioning.

Accurate Positioning





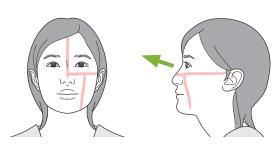
Patient Facing Down





V-shaped Dental Arch

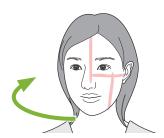
Patient Facing Up

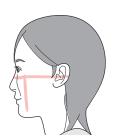


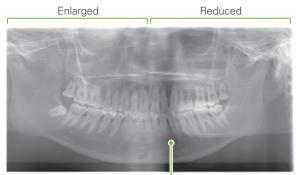


Jaw joint is out of picture

Patient Facing to the Right -

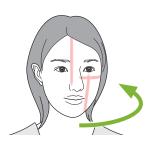


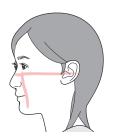


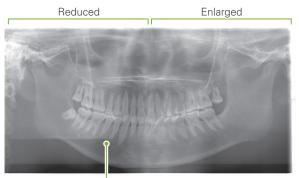


Reduced X-ray absorption here (excessive saturation)

Patient Facing to the Left

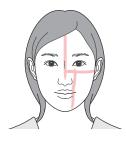


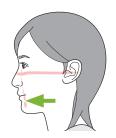




Reduced X-ray absorption here (excessive saturation)

● Front-Back Beam is too far forward than the distal side of the left canine teeth.

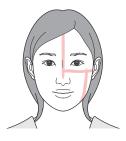


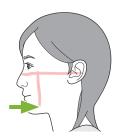




Enlarged

● Front-Back Beam is too far back than the distal side of the left canine teeth.



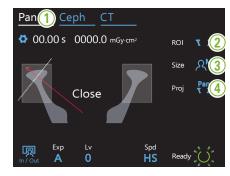




Reduced

6.4 Quadruple TMJ Exposure

6.4.1 Preparation



1. Select panoramic exposure mode

Touch the Pan key to set the unit for panoramic exposure mode.

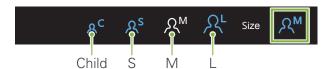
2. Select region of interest (ROI)

Touch the icon to the right of "ROI" to select the TMJ.



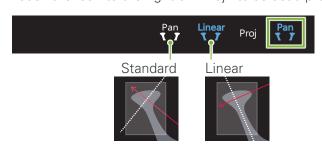
3. Select patient size

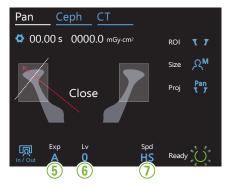
Touch the icon to the right of "Size" to select the patient size.

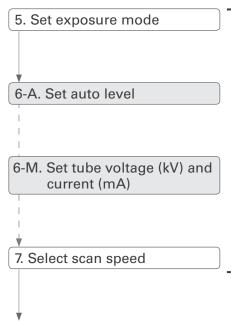


4. Select projection

Touch the icon to the right of "Proj" to select a projection.

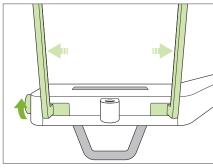


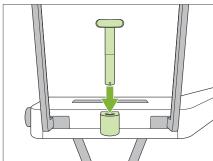


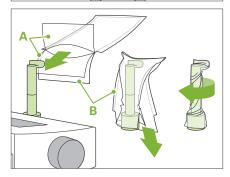


Same as for dental arch panorama. Refer to 6.3.1.1,

"7. Set exposure mode" – "9. Select scan speed" (pp. 27–28).







A: Cover (semi-transparent)

B: Paper (white)

8. Set upper lip rest

Turn the temporal stabilizer knob and open the stabilizers.

Wipe the upper lip rest with Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) and put it in the chinrest holder.

* Before use, make sure the upper lip rest is not scratched or damaged in any way.

Put a chinrest disposable cover on the upper lip rest.

- 1. Open the cover and put it on the upper lip rest.
- 2. Pull down the paper and peel it off.
- 3. Twist the cover and press it against the upper lip rest as tightly as possible.

⚠WARNING

- The upper lip rest must be disinfected by wiping the Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) after each use, and then put a chinrest disposable cover on the upper lip rest.
- Chinrest disposable covers must never be re-used. A new one must be used for each patient.
- The patient handle must be disinfected by wiping it with Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) after each patient. If the patient has an open or bleeding wound on his hand, the patient handle should be covered to prevent cross contamination before the patient grips it.

Store chinrest disposable covers in a clean, hygienic place.

6.4.2 Patient Entry and Positioning

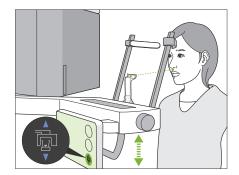
For patient positioning, refer to 2 Precautions, "Patient Size and Positioning" (p. 9).

1. Prepare patient

Have the patient wear an X-ray protection apron.

ACAUTION

- The patient must wear an X-ray protection apron.
- Patient must remove glasses and all accessories like necklaces etc. Otherwise the exposure could fail.
- Make sure the patient's hair cannot get caught in moving parts. If the patient's hair is tied in the back like a pony tail etc., have the patient undo it. Otherwise, the arm might hit the patient.



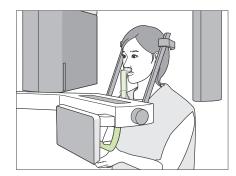
2. Adjust unit height

Adjust the height of the unit so that the upper lip rest is even with the patient's upper lip.

Press and hold the Up and Down switches to raise or lower the unit; release the switch to stop the unit.

ACAUTION

- Take care that the temporal stabilizers do not strike the patient in the eye.
- Take care that the patient is not hit or pinched by any part of the lift assembly (control panes support, arm, arm support).
- Always use the Up and Down switches to adjust the unit's height. Never try to force it; this would damage the unit.
- If the lift reaches its upper limit, a series of beeps will sound. If at this point the Ready key is pressed, a message will appear saying that this is the limit for a CT exposure. If you make a CT at this point, the maxillary region may not be fully exposed. In this case, have the patient sit down or lower his position in some other way.



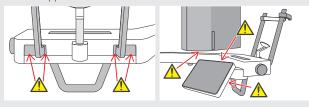
Positioning Point Correct Posture

3. Patient entry

Have the patient move forward and put his upper lip on the upper lip rest. Have the patient lightly hook his thumbs on the patient handle.

ACAUTION

- Do not exert force to have the patient enter or leave; this could damage the temporal stabilizers.
- Never let the patient touch any of the control panel keys and switches.
- Keep fingers away from gaps and openings of moving parts as well as the holes on the support column.



• The upper lip rest can withstand a load of 2 kg and the patient handle and control panel support a load of 5 kg. These parts could be broken or the patient injured if the patient rests too much body weight on them.

Patient stands straight and neck is as perpendicular as possible.

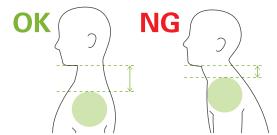
If the patient stands (or sits) too far back, the upper torso will lean forward and the neck will be slanted. Have the patient move forward in this case.

Pull jaw in and make the line from the orbitale to the ear orifice horizontal.

To get correct posture, it helps to tell the patient to move his forehead forwards while keeping his neck stretched while the lift is being lowered.

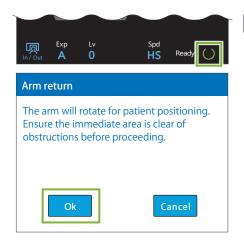
Relax and lower shoulders.

If the shoulders are too close to the jaw, the arm might hit them. For patients with square shoulders or a short neck, have them relax and lower their shoulders as much as possible. If the neck is slanted, have the patient straighten it and pull in his jaw. If the shoulders are too tension, the patient may be gripping the patient handle too tightly. Have patients lightly hook their thumbs on the handle instead.



CAUTION

• For patients who have square shoulders or short necks, the arm might contact the shoulder. In this case, have the patient lower his shoulders as much as possible and then run a test with no X-ray mode to make sure the arm does not contact the patient.



4. Touch Ready key

Touch the Ready key.

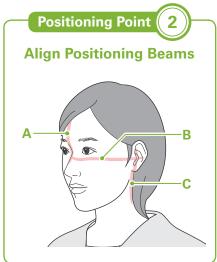
The "Arm return" dialogue will appear. Make sure the area is safe, and then touch the "Ok" button.

① If a dialogue box asking to check the temporal stabilizers appears, touch the "Ok" button.

The arm will move to its patient position and the mid-sagittal, horizontal and front-back beams will light up.

MWARNING

•The laser beam could damage a person's eyesight; never look directly into the beam and let it strike anyone's eyes.





Front-Back Beam Switches
Forward

Backward

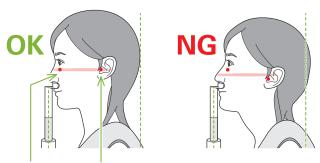
5. Line up beams

A: Mid-sagittal beam lines up with mid-sagittal plane.

Move the patient's head to line it up with the beam.

B: Horizontal beam lines up with the line from orbitale to ear orifice.

Have the patient keep a posture with the line from orbitale to ear orifice horizontal and then press the Up or Down switches to line the beam up with it.



Beam passes through these two points.

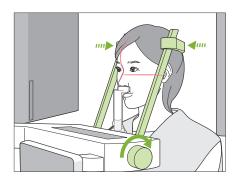
ACAUTION

• If the line from the orbitale to the ear orifice is not horizontal, the back of the head might stick out and make contact with the arm during an exposure or ready move. For a patient who has a large head, run a test with no X-ray emission mode to make sure the arm does not contact the patient.

C: Front-back beam lines up with the outer ear orifice.

Press the Forward or Backward switches to move the beam so it is properly lined up.

* The image layer is about 12 mm in front of the front-back beam.



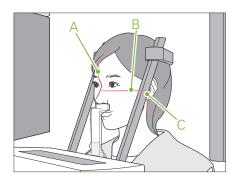
6. Close temporal stabilizers

Adjust the height of the temporal stabilizers.

Turn the temporal stabilizer knob and close the stabilizers firmly. Have the patient close his mouth.

ACAUTION

• Do not use excessive force to close the temporal stabilizers. This could be uncomfortable for the patient or damage the stabilizers.



7. Check beams

Check the positioning of the all beams.

Mid-sagittal beam (A) lines up with mid-sagittal plane.

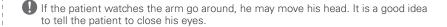
Horizontal beam (B) lines up with the line from orbitale to ear orifice.

Front-back beam (C) lines up with the outer ear orifice.

Tell the patient not to move his head during X-ray emission (or while the melody is playing) and then leave the X-ray room.

▲CAUTION

• Warn the patient not to move during emission (while the melody is sounding). If the patient moves, he could be hit by the arm or the image might be ruined.



Min. Scan Move Function

(If this function is turned on.)

Before leaving the X-ray room, the arm can be moved to the position which will require the shortest exposure time (least time needed to hold down the Emission switch).

8. Touch Ready key

Make sure the patient has his hands on the patient handle. With the unit already in the ready state, press the Ready key again. The arm will move to its Min. Scan Move position and then a two-toned beep will sound.

Min. Scan Move:

- X-ray emission will begin as soon as the Emission Switch is pressed.
- Only the Ready key will work.
- * Touch the Ready key again the arm will move back to its original patient positioning position.
- ① If after the arm moves to the Min. Scan Move position, one of the beams is not properly line up, touch the Ready key again to move the arm back to its original patient positioning position and re-position the patient.

After the arm moves, re-check the patient's positioning and hand position on the patient handle and then leave the X-ray booth.

6.4.3 Exposure

1. Check ready state

Check that the Ready key on the control panel and the Ready LED (green) on the control box are on.



2. First X-ray emission (mouth closed)

Hold down the Emission switch.

The arm will start to rotate and exposures will be made of the left and right sides.

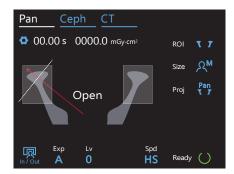
The Emission LED (yellow) on the control box will light up and the melody will sound.

▲WARNING

• If an emergency occurs, release the Emission switch. Or press the Emergency Stop switch. This will stop the arm rotation and X-ray emission.

ACAUTION

- Go outside the X-ray room to press the Emission switch.
- If the operator must stay in the X-ray room for some reason, he should wear an X-ray protection apron and stay at least 2 meters away from the X-ray emission focal point. Also he should stay outside the area exposed to X-ray emission.
- Releasing the Emission switch during X-ray emission will stop the arm's rotation and terminate the exposure. In this case guide the patient away from the unit. If the X-ray head or the detector is directly in back of the patient, turn off the unit and slowly rotate the arm so the patient can leave. If you perform the ready procedures without having the patient move away from the unit, the arm could hit and possibly injure the patient.
- If an error occurs during an exposure, the arm will stop and the exposure will be terminated. In this case guide the patient away from the unit. If the X-ray head or the detector is directly in back of the patient, turn off the unit and slowly rotate the arm so the patient can leave. If you perform the ready procedures without having the patient move away from the unit, the arm could hit and possibly injure the patient.
- ① As much as 15 seconds could elapse from the time the Emission switch is pressed until X-ray emission actually starts. This is not abnormal. It takes some time for the unit to verify the computer settings.
- If the computer is not ready, an error message will appear in the control panel. If this happens, turn off the Power switch. When the computer is ready, turn on the unit again.
- * After the first exposure, the message "Exposure Standby" will appear in i-Dixel. If you select another mode at this time, the image for the first exposure will be transmitted to the computer.



3. Check ready state

When the melody stops, release the Emission switch and hang it in its holder on the control box and go inside the X-ray room.

Have the patient maintain the proper posture and touch the Ready key. The arm will go back to its starting position.

"Open" will appear in the control panel. Have the patient open his mouth.

4. Second X-ray emission (mouth opened)

This is done in the same way as the 1st emission.



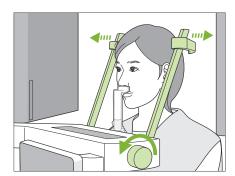
5. Exposure completed

The melody stops when the exposure has been completed.

The arm will automatically move to the patient egress position.

Release the Emission Switch and hang it in its holder on the control box.

6.4.4 Patient Egress



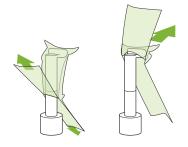
1. Open temporal stabilizer

Turn the temporal stabilizer knob and open the stabilizers completely.

Guide the patient away from the unit.

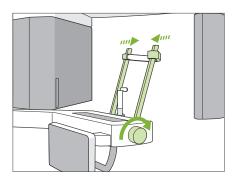
ACAUTION

- Take care that the temporal stabilizers do not strike the patient in the eye.
- Be careful of the stabilizers when the patient leaves the unit.



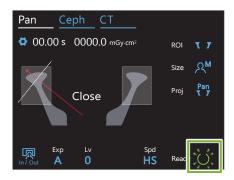
2. Throw away disposable covers etc.

Throw away the chinrest disposable cover on the upper lip rest. If the upper lip rest cover is removed by splitting its seams and turning it inside out, the outer surface will not touch anything.



3. Close temporal stabilizer

Turn the temporal stabilizer knob and close the stabilizers completely.



4. Touch Ready key

Touch the Ready key.

The "Arm return" dialogue will appear. Make sure the area is safe, and then touch the "Ok" button.

The arm will move back to its original patient positioning position.

* If another exposure will be made right away, touch the In/Out key to move the arm to its patient entry position.

6.4.5 Image Transmission

Same as for dental arch panoramic exposure.

Refer to "6.3.5 ImageTransmission" (p. 40).

7 CT Exposure

7.1 Exposure Types and Functions

7.1.1 FOV (Field of View)

There are 8 types of exposures for varying diagnostic purposes.

Ø40

For 2 or 3 teeth or the jaw joint. X-ray dosage is very low because exposure is localized.

Ø80

For the entire dental arch.

Wisdom teeth of the back molar might not get into the image.

* R100 and F150 only.

■ Ø100 (dental arch FOV)

MORITA's unique FOV that fits the dental arch. An image of the entire dental arch is made with a lower X-ray dosage.

- * Ø100 equivalent.
- * For patients with large jaws, the whole region might not be in the exposure.
- * R100 and F150 only.

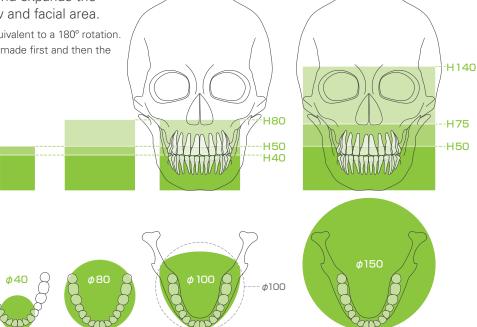


Ø100 × H80

Ø150

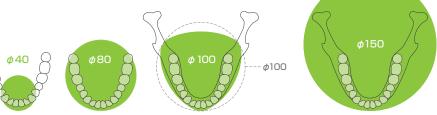
This exposures the entire jaw area. The H140 exposure combines two exposures and expands the exposure range to the entire jaw and facial area.

- * The arm rotates 360°, but the data is equivalent to a 180° rotation.
- * For an H140 exposure, the lower part is made first and then the upper part.
- * F150 only.



Exposure Region Dimensions

* Depending on the patient's body size and shape of the dental arch, actual exposure range might not fit as the illustration.



Exposure Functions Chart

Field of View (FOV)	Scan	(Scan)	Resolution (Res)			
	180°	360°	HR (high resolution)	SD (standard)		
Ø40 × H40	0	0	0	0		
Ø40 × H80	0	0	-	0		
Ø80 × H40 Ø80 × H50 Ø80 × H80	0	0	-	0		
*Ø100 × H40 *Ø100 × H50 *Ø100 × H80	0	-	-	0		
Ø150 × H50 Ø150 × H75 Ø150 × H140	0	-	-	0		

Order Selection							
F40	R100	F150					
0	0	0					
0	0	0					
-	0	0					
-	0	0					
-	-	0					

^{*} Ø100 equivalent.

7.1.2 180° & 360° Exposure Modes

Data for image reconstruction can be obtained with either 180° or 360° scans.

180°

Use this for patients who find it hard to stand still or to reduce X-ray dosage.

360°

Since the data obtained has a higher density, there are fewer artifacts in the image.

- * For a CT exposure of a large or larger than permitted patient, use 180° exposure mode to keep the arm from contacting the patient.
- * For the Ø150, the arm rotates 360°, but the data is equivalent to a 180° rotation.

180° Exposure



360° Exposure



7.1.3 Resolution (Res)

■ High Resolution (HR)

High resolution images are ideal for treatment of periapical lesions, periodontics, etc. Voxel size is 80 µm.

Patient Size	2.5 lp/mm MTF (%)		
C (Children), S, M	10		
L	7		



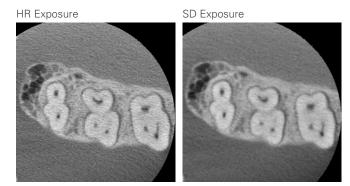


- * Even slight patient movement can greatly affect the image quality. Consider doing the following things to make sure the patient stays as still as possible.
- Tell the patient to close his eyes during the exposure.
- For greater stability, it is better to have the patient sit on a chair.
- Using the 180° exposure is recommended if it is hard for the patient to stay still for a long time.



If you wish to acquire better images, select "M" at the exposure settings. Selecting DR (dose reduction mode) emits less X-rays but images are grainier.

* For details, refer to 7.3.1, "7. Set exposure mode" (p. 65).



i-Dixel Operation and Resolution

Zoom Reconstruction Function

A higher resolution image (80 μ m pixels) can be made without repeating the exposure by specifying a region of a completed CT image and having that area reconstructed again.

Slice Thickness Adjustment

After making a CT exposure, the slice thickness can be adjusted with i-Dixel. Adjusting the slice thickness has the following effects:

- Thicker slices have lower resolution but make smoother images.
- Thinner slices have higher resolution but are grainier.

0.96 mm Slice (standard setting)



Slice Thickness: 0.48 mm

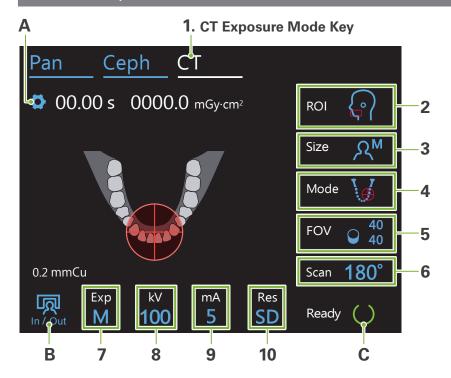


Since the pixel size for high resolution (HR) is 80 μ m, a 0.48 mm slice should produce better resolution than the standard resolution (SD). Adjust the slice thickness to suit your purposes such as observing delicate features of hard tissue.

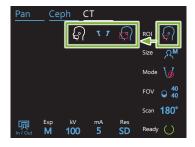
* For details refer to the Operation Instructions for i-Dixel.

7.2 Operation and General Settings

7.2.1 CT Exposure



Numbers 2 to 10 show the current settings. Touch any one of the icons to show other selections.





X-ray Dosage

A. Settings Key

Hold down this key to save the current settings.

B. In/Out Key

Touch for patient entry and egress. The arm moves to 90° to the lift for easy patient entry and egress.

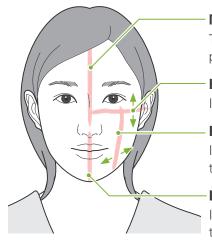
C. Ready Key

Shows that the unit is in the ready state.

This key blinks when the Power switch is turned on.

Touch it after guiding the patient into position: it will light up. The arm will move to its patient position and the mid-sagittal, horizontal, and front-back (left-right) beams will light up. The unit is ready to begin X-ray exposure whenever the Emission switch is pressed.

7.2.2 Positioning Beams



Mid-Sagittal Beam

This beam lines up with the patient's mid-sagittal plane. Press the Beam On/Off switch to turn it on.

Horizontal Beam

Indicates the height center of the FOV.

Front-Back Beam **◄** ▶

Indicates the center in the front-back direction of the FOV.

Left-Right Beam <

Indicates the center in the horizontal direction of the FOV.

Beam On/Off Switch (Mid-sagittal, Horizontal, Front-Back)



Beam On/Off Switch (Left-Right)



7.2.3 Settings

2. Exposure Area (ROI)	3. Patient Size (Size)	4. Positioning Modes (Mode)	5. Exposure Region (FOV)		6. Scanning (Scan)	7. Exposure Setting (Exp)	8. Tube Voltage (kV)	9. Tube Current (mA)	10 Resolution (Res)	Rest Method		
(1101)	(0120)	(111040)	Ø	Н		(EXP)	(KV)	(11174)				
		Panoramic Scout (for Ø40 FOV only)	40	40	180° (Half Scan)	DR (Dose Reduction)	100		HR (High Resolution) SD (Standard)	- Chinrest		
		Two-Direction Scout		80				2-8		Cillinest		
Dental Arch	AC	<u>i i i i i i i i i i </u>		40		M						
$\left\{ \cdot \right\}$		****	80 ^{* 1}	50	360° (Full Scan)	(Manual)	95			Bite Block		
				80	(i dii Oddii)				SD (Standard)			
		Beam Positioning		40		Off	90			DITC DIOCK		
٨s		100 [*] 1	50	180° (Half Scan)	(No X-ray		2-9					
		'		80		Emission)		2-3				
TMJ	$\mathcal{L}_{\mathbf{M}}$	Two-Direction Scout		40	180° (Half Scan)		85	85		35]	HR (High Resolution) SD (Standard)	Upper Lip
7 7	Beam Positioning	40	80	360° (Full Scan)	M (Manual)	80 7			Rest			
Maxillofacial	\mathcal{N}_{Γ}			50		Off (No X-ray	75	2 – 10	SD (Standard)	Chinrest		
		Beam Positioning	150 ^{*2}	75		Emission)	70 _					
		"		140						Bite Block		

*1 R100 and F150 only. *2 F150 only.

■ Patient Size and Tube Voltage / Current



Tube voltage and current are adjusted according to patient size.

The set values for patient size is the sagittal length of the skull (A).

Also the upper limit for the lift cannot accommodate patients taller than 180 cm. We recommend having patients who are too tall to sit down.

	C (Child)	S (Small)	M (Middle)	L (Large)
Sagittal Length (A)	Max. 17 cm	Max. 17 cm	Max. 19cm	Max. 21cm
Tube Voltage	100 kV	100 kV	100 kV	100 kV
Tube Current	3 mA	3 mA	5 mA	7 mA

^{*} The above values are estimates only. The dentist should judge this by the patient's body shape, skeletal structure etc.

Positioning Modes

There are three ways to perform positioning to make a CT exposure: the panoramic scout, the two-direction scout, and beam positioning.



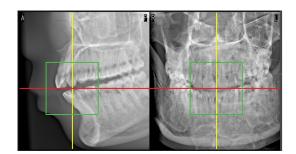
Panoramic Scout (i-Dixel)

Determine the exposure area (FOV) using a panoramic image. A panoramic image previously made with the unit may be used.

* Depending on the patient, there may be a large error in the positioning for mandibular second and third molars. In this case, it is better to use the two-direction scout

Intended ROI: Dental Arch

Intended FOV: Ø40



Two-Direction Scout

Determine the exposure area using two images, one lateral and the other posterior-anterior.

Intended ROI: Dental Arch, TMJ

Intended FOV:

Dental Arch: Ø40, Ø80, Ø100

TMJ: Ø40



Beam Positioning

Determine the exposure area by lining up the beams with the patient without the aid of i-Dixel.

* For edentulous patients, it is hard for scout positioning to be reproduced for a CT scan, and beam positioning should be used instead.

Intended ROI: All Intended FOV: All

Rest Method



Chinrest

Use this to make an exposure of natural occlusion or when the bite-block cannot be used for edentulous patients or some other reason.



Bite-block

Since this lines up with the upper and lower incisors, there is a high degree of reproducibility.

* Part of the bite-block will appear in the image.



Upper Lip Rest

This is mainly used for exposures of the TMJ.

7.3 CT Exposure

7.3.1 Preparation



1. Select CT mode

Touch the CT key to set the unit for CT exposure mode.

2. Set exposure area

Touch the icon to the right of "ROI" to select the region of interest.



3. Select patient size

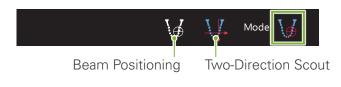
Touch the icon to the right of "Size" to select the patient size.



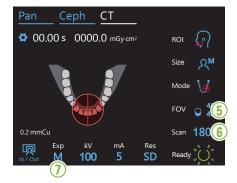
4. Check positioning mode

Make sure the "Mode" is set for "Beam Positioning".

- * For using the panoramic scout, this is not necessary. Display the panoramic image to be used in i-Dixel.
- * Even for a two-direction scout exposure, patient positioning is performed by the beam positioning mode.



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5. Select exposure area (FOV)

Touch the icon to the right of "FOV" to select exposure area.

Dental Arch CT

(R100)



① For Ø100 exposures, the arm may contact the patient's shoulder depending on the patient's body shape. The probability of this happening can be reduced by using the Ø80 FOV instead.

(F40)



TMJ CT



Maxillofacial CT * F150 only.



6. Select scan mode

* Ø40 and Ø80 only.

Touch the number to the right of "Scan" to select scan mode.

- **180°**: The arm rotates for half the circumference around the head (half-scan). This reduces X-ray dosage.
- **360°**: This is a full scan where the arm rotates all the way around the head.

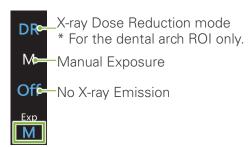
This provides better image quality and reduces artifacts.



* For a CT exposure of a large or larger than permitted patient, use 180° exposure mode to keep the arm from contacting the patient.

7. Set exposure mode

Touch the letter under "Exp" to set the exposure mode.



DR (Dose Reduction mode)

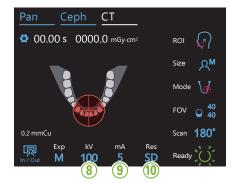
- •X-ray dosage is reduced by reducing X-ray emission for relatively transparent areas and increase it for more opaque areas. This also improves the delineation of the outlines of areas where there are hollows such as the checks and jaws. X-ray dosage will be reduced by as much as 60% compared to the standard exposures. * The amount of dosage reduction depends on the exposure mode setting.
- "DR" cannot be selected for TMJ exposures.
- "DR" cannot be selected for the two-direction scout exposures.
- •When "DR" has been selected, "Max" will appear next to the tube current value. This means the displayed value is the maximum mA output, although it will vary during the scan. In DR mode, tube current can be set from 3 to 10 mA. (From 3 to 8 mA if tube voltage is 85 kV or more.)



• "DR" will only work if tube current is set for 3 mA or more. In DR mode, if the mA is set for less than 2 mA, it will automatically go up to 3 mA when the unit is set for DR mode.

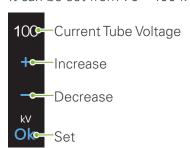
Exposure Settings for High Resolution

To get the full benefit of the high resolution "HR" setting, use the manual "M" exposure settings. Selecting "DR" (dose reduction mode) emits less X-rays but images are grainier.



8. Set tube voltage (kV)

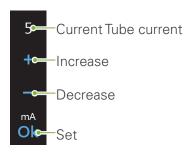
Touch the number under "kV" to set the voltage. It can be set from 70 – 100 kV in 5 kV increments.



Patient Size			S	М	L
Estimated Tube Voltage (kV)	CT Exposure	100	100	100	100
	Panoramic/ Two-Direction Scout Exposure	90	90	90	90

9. Set tube current

Touch the number under "mA" to set the tube current. It can be set from 2 – 10 mA in 1 mA increments.



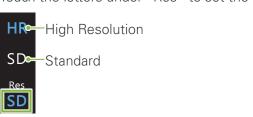
Patient Size			S	M	L
Estimated Tube Current (mA)	CT Exposure	3	3	5	7
	Panoramic/ Two-Direction Scout Exposure	2	2	2	2

^CAUTION

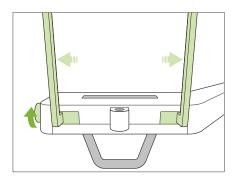
• Strong X-ray emission can saturate the photo diodes on the sensor for areas having high X-ray transparency. In the resulting image, this might falsely appear to be a complete absence of tissue. Watch the rotating image produced during -ray emission to see if this happens and take this into consideration when making a diagnosis.

10. Select resolution $* \emptyset 40 \times H40$ only.

Touch the letters under "Res" to set the resolution.







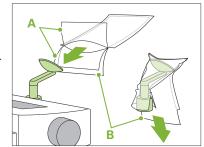
Turn the temporal stabilizer knob and open the stabilizers.

Wipe the chinrest, bite-block, or upper lip rest with Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) and put it in the chinrest holder.

* Before use, make sure the component is not scratched or damaged in any way.

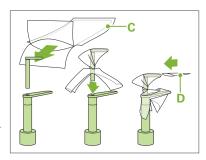
Chinrest

- 1. Open a chinrest (A) disposable cover and put it on the chinrest.
- 2. Pull down the paper (B) and peel it.



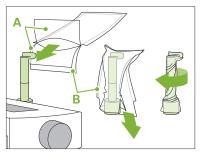
Bite-Block

- 1. Peel off the paper of a chinrest disposable cover (C) and cover the bit (the part of the bite-block the patient bites on) with it.
- 2. Twist the disposable cover and put the bit in the bite-block stand.
- 3. Put a bite-block cover (D) on the bit.



Upper Lip Rest

- Open the chinrest disposable cover (A) and put it on the upper lip rest.
- 2. Pull down the paper (B) and peel it.
- 3. Twist the cover and press it against the upper lip rest as tightly as possible.



WARNING

- •The chinrest, bite-block, and upper lip rest must be disinfected by wiping with Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) after each use, and before putting on a disposable cover.
- Disposable covers for the chinrest and bite-block must never be re-used. A new one must be used for each patient.
- The patient handle must be disinfected by wiping it with Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) after each patient. If the patient has an open or bleeding wound on his hand, the patient handle should be covered to prevent cross contamination before the patient grips it.

ACAUTION

- The chinrest, bite-block and upper lip rest can withstand a load of 2 kg. These parts could be broken or the patient injured if the patient rests too much body weight on them.
- Store disposable covers for the chinrest and bite-block covers in a clean, hygienic place.

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7.3.2 Patient Entry and Positioning



Dental Arch CT Exposure

Same as for dental arch panoramic exposure.

Refer to "6.3.2 Patient Entry and Positioning" (pp. 34–37).

For Panoramic Scout:

Press the Pan key on the control panel to go to panorama mode and then position the patient. If the patient is positioned for any mode other than panorama, the scout position may not be accurate.

If you use a panoramic image made previously, make sure patient positioning is exactly the same.



TMJ CT Exposure

Same as for quadruple TMJ exposure.

Refer to "6.4.2 Patient Entry and Positioning" (pp. 47-51).



Maxillofacial CT Exposure

Same as for dental arch panoramic exposure.

Refer to "6.3.2 Patient Entry and Positioning" (pp. 34-37).

① Since there is an upper limit for the exposure, it may be necessary to have very tall patients sit down or lower their posture.

7.3.3.1 Set FOV Position with Panoramic Scout (for Ø40 FOV)

CAUTION

- The panoramic scout determines a reference range, but its accuracy cannot be guaranteed. Depending on the patient, there may be a large error in the positioning for mandibular second and third molars. In this case, it is better to use the two-direction scout.
- For edentulous patients, it is hard for scout positioning to be reproduced for a CT scan. In this case, use beam positioning to make the CT exposure.



1. Display a panoramic image (i-Dixel)

Display a panoramic image in i-Dixel.

To make a new panoramic image refer to "6.3.3 Exposure" (p. 38).

The panoramic scout can be used with panoramic images made with the Veraviewepocs 2D, 3D, and 3D R100/F40 models. However, since the positioning will be lower, always check it with the beams before making the CT exposure.

ACAUTION

• If the panorama is to be used only for CT exposure positioning, use the lowest X-ray dosage possible without having an adverse effect on CT positioning. Estimated Tube Voltage and Current: 90 kV and 2 mA for all patient sizes.



2. Open X800 Exposure window

Click the Panoramic Scout Image Select icon (A).

When the panoramic image opens in i-Dixel, the panoramic scout procedure can be made.



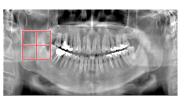
A green frame will appear. This frame specifies the exposure area (FOV).



3. Specify exposure area (i-Dixel)

Drag the frame to determine the center of the exposure.

- Repeated clicking or dragging for a long time can slow down the computer's reaction or even freeze it.
- ① Do not touch any operation keys on the X-ray unit while the exposure area frame is being displayed in i-Dixel. This could cause the X-ray unit to stop working.
- If the frame goes out of the possible range of exposure, it will turn red, and it will not be possible to specify the center of the FOV.



Changing the FOV on i-Dixel

The FOV can be changed with i-Dixel.

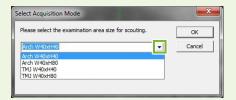
Click the Set CT Area icon

Click the Set CT Area icon (A) on the scout panel to display a exposure mode selection dialogue box.



Select area size

Select an area size from the pull-down menu and then click the "Ok" button.



* If positioning mode is Two-Direction Scout for TMJ, the exposure will be "Size L" regardless of the selected patient size due to the limitation of arm movement. Select appropriate area size from any of the followings:

W 40 × H 80 (Size L)

W 40 × H 40 (Size L)

W 40 × H 40 (HR Size L)

If other is selected, the frame will turn red, and it will not be possible to make an exposure.

The area size will change

A beep will sound and the area size will change to the one selected.





Scout position check

Press OK to move the unit to the scouted position. The arm will move in multiple directions so ensure the immediate area is clear.

Press Cancel to start the scout operation from the beginning.





4. Send FOV position (i-Dixel)

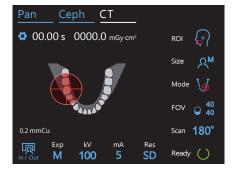
Click the Send CT Scout Position icon (A) on the scout tool panel.

A yellow cross mark "+" in the panoramic image indicates the center of the exposure area.

① Do not fail to click the Send CT Scout Position icon after positioning the frame. Otherwise, the newly selected area will not be recognized by the Veraview X800.

If the selected position is OK, a two-toned beep will sound and the message shown to the left will appear in the control panel. Touch the "Ok" button to confirm the positioning.

Touch the "Cancel" button to cancel it.



5. Register FOV position

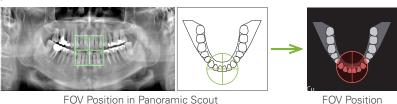
Touch the "Ok" button, then the arm will move into position correspond to the registered FOV position.

The control panel will automatically change to CT exposure mode. The exposure area specified by i-Dixel will appear as an FOV circle in the control panel.

The FOV circle determines a reference range, but its accuracy cannot be guaranteed.

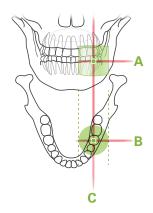
Incisal Positioning

When the scout indicates the incisal region, the FOV circle in the control panel may appear slightly back. This is because the unit has adjusted the position so that the FOV is within the exposure range of the unit. The incisal region will not be exactly in the center of the FOV, but it will be included in the exposure range. Continue with the procedure.



Determined by the Veraview X800

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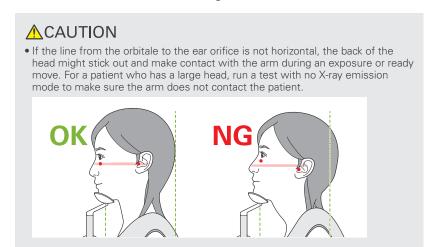


Front-Back Beam (B) Switches
Forward Backward

6. Check exposure position

The horizontal (A), front-back (B), and left-right (C) beams will intersect at the center of the exposure area.

Check that the beams are on target.



Tell the patient not to move his head during X-ray emission (or while the melody is playing) and then leave the X-ray room.



• Warn the patient not to move during emission (while the melody is sounding). If the patient moves, he could be hit by the arm or the image might be ruined.

If the patient watches the arm go around, he may move his head. It is a good idea to tell the patient to close his eyes.

Min. Scan Move Function

(If this function is turned on.)

7. Touch Ready key

Make sure the patient has his hands on the patient handle. With the unit already in the ready state, press the Ready key again.

The arm will move to its Min. Scan Move position and then a two-toned beep will sound.

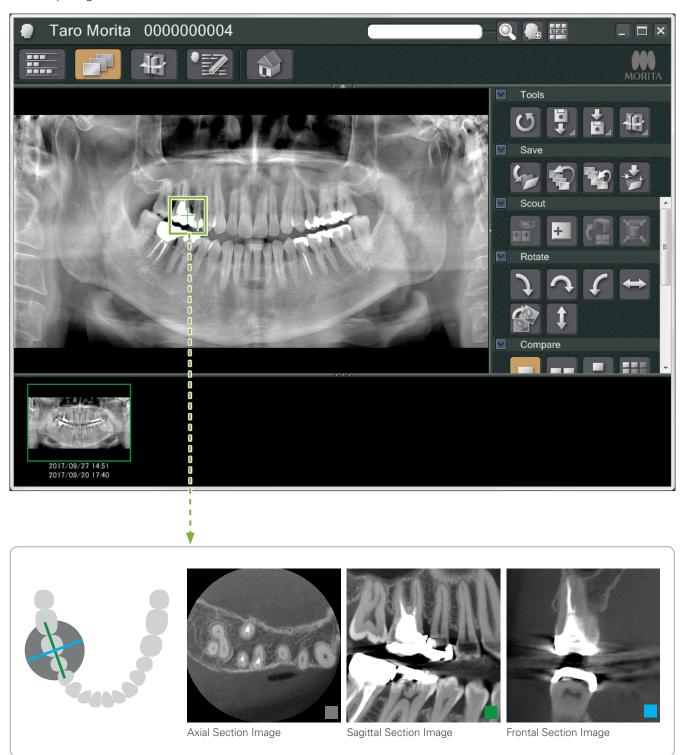
After the arm moves, re-check the patient's positioning and hand position on the patient handle and then leave the X-ray room.

To continue go to "7.3.4 CT Exposure" (p. 81).

Relationship between CT and Panoramic Scout Images

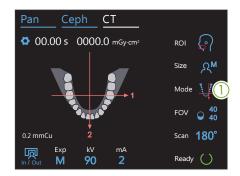
The CT and panoramic scout images are linked. If you double-click the green cross mark "+" in the panoramic scout image, the CT image for that place will be automatically displayed.

If the CT exposure has not been made, the cross mark "+" will be yellow. In this case, there is no linkage with anything.



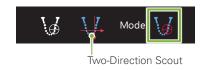
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7.3.3.2 Set FOV Position with Two-Direction Scout (for Dental Arch and TMJ ROIs)



1. Select two-direction scout

Touch the icon to the right of "Mode" to select "Two-Direction Scout". The positioning beams will turn off.





Arm return

The arm will rotate for patient positioning. Ensure the immediate area is clear of obstructions before proceeding.



Cancel

2. Touch Ready key

Touch the Ready key.

The "Arm return" dialogue will appear. Make sure the area is safe, and then touch the "Ok" button.

If a

If a dialogue box asking to check the temporal stabilizers appears, touch the "Ok" button.

The X-ray head will move and the positioning beams (horizontal, front-back, and left-right) will light up. The unit is now in the ready state. The horizontal beam moves to the scout position.

MARNING

• The laser beam could damage a person's eyesight; never look directly into the beam and let it strike anyone's eyes.



A Left-Right Beam Switches

Left

Right

B Front-Back Beam Switches

Forward

Backward

3. Line up beams

Make sure the FOV positioning: "Mode" is set for two-direction scout. Line up the front-back and left-right beams with the center of the target area.

* The exposure range for the two-direction scout is W80 \times H80 (W40 \times H80 for F40 models). If these beams are too far off the target, it may not be contained within the two-direction scout image.

Reference Point for Exposure Position

Dental Arch

Front-Back Beam: first molar

TM.

Front-Back Beam: 1 cm in front of ear orifice

Left-Right Beam: seen head-on, patient's TMJ position

For a TMJ exposure, make sure the "ROI" is set for "TMJ". The scout and CT exposure ranges for the dental arch and the TMJ are not the same.



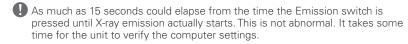
4. Check ready state

Check that the Ready key on the control panel and the Ready LED (green) on the control box are on.



5. Execute two-direction scout exposure

Hold down the Emission switch. The arm will rotate and X-ray emission will start. The Emission LED (yellow) on the control box will light up and the melody will sound.

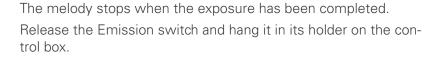


MWARNING

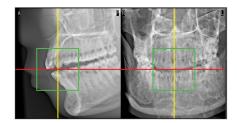
• If an emergency occurs, release the Emission switch. Or press the Emergency Stop switch. This will stop the arm rotation and X-ray emission.

ACAUTION

- Go outside the X-ray room to press the Emission switch.
- If the operator must stay in the X-ray room for some reason, he should wear an X-ray protection apron and stay at least 2 meters away from the X-ray emission focal point. Also he should stay outside the area exposed to X-ray emission.
- If the Emission switch is released before completing the entire scout exposure process, the exposure will be immediately terminated. In this case guide the patient away from the unit. If the X-ray head or the detector is directly in back of the patient, turn off the unit and slowly rotate the arm so the patient can leave. If you perform the ready procedures without having the patient move away from the unit, the arm could hit and possibly injure the patient.
- For the two-direction scout, hold the Emission switch down until both exposures are completed. Releasing it after only one has been made will result in losing the first exposure and not getting any image at all.







6. Scout image display (i-Dixel)

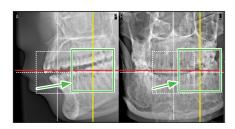
The two-direction scout image will be displayed in the capture window.

The intersection of the red and yellow lines shows the center of the exposure area and the green frame shows the size of the FOV

The size of the frame reflects the size of the selected FOV.

ACAUTION

- Do not close the scout image until the CT exposure has been completed. If the scout image is closed, positioning will no longer be possible.
- * The CT exposure area size (FOV) can be changed with i-Dixel. For details, refer to 7.3.3.1, "Changing the FOV on i-Dixel" (p. 70).

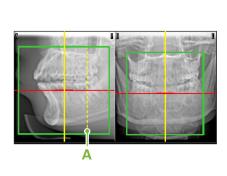


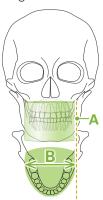
7. Specify FOV position (i-Dixel)

Drag the frame to determine the center of the FOV.

● Ø100 Scout Image * R100 only.

A dotted yellow line (A) appears in the image for the Ø100 FOV. This line represents the maximum width position of the FOV (B). Use this line to estimate the positioning.

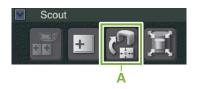




- Repeated clicking or dragging for a long time can slow down the computer's reaction or even freeze it.
- ① Do not touch any operation keys on the X-ray unit while the exposure area frame is being displayed in i-Dixel. This could cause the X-ray unit to stop working.
- If the frame goes out of the possible range of CT exposure, it will turn red, and it will not be possible to specify the center of the FOV.



76



Press OK to move the unit to the scouted position. The arm will move in multiple directions so ensure the immediate area is clear. Press Cancel to start the scout operation from the beginning. Ok Cancel

Ok Cancel Pan Ceph CT

8. Send FOV position (i-Dixel)

Click the Send CT Scout Position icon (A) on the scout tool panel. A cross mark "+" will appear at the center of the exposure area in the image.

① Do not fail to click the Send CT Scout Position icon after positioning the frame. Otherwise, the newly selected area will not be recognized by the Veraview X800.

If the selected position is OK, a two-toned beep will sound and the message shown to the left will appear in the control panel.

Touch the "Ok" button to confirm the positioning.

Touch the Cancel button to cancel it.



9. Register FOV position

Touch the "Ok" button, then the arm will move into position correspond to the registered FOV position.

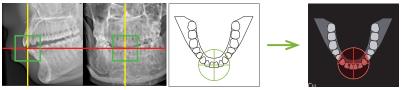
The exposure area specified by the two-direction scout will appear as an FOV circle in the control panel.

1 The FOV circle determines a reference range, but its accuracy cannot be guaranteed.

Incisal Positioning

When the scout indicates the incisal region, the FOV circle in the control panel may appear slightly back. This is because the unit has adjusted the position so that the FOV is within the exposure range of the unit. The incisal region will not be exactly in the center of the FOV, but it will be included in the exposure range. Continue with the procedure.

* Note that the FOV position specified on the scout image differs from the position displaying on the control panel.



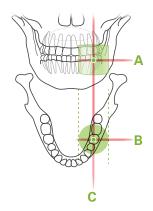
FOV Position on Two-Direction Scout

FOV Position Determined by the Veraview X800

10. Set exposure mode

Set the exposure (Exp), tube voltage (kV), and current (mA) settings for the CT exposure.

For details, refer to 7.3.1, "7. Set exposure mode" – "9. Set tube current" (pp. 65–66).





Left-Right Beam (C) Switches





Horizontal Beam (A) Switches

▲Up ▼Down

Front-Back Beam (B) Switches

Forward Backward

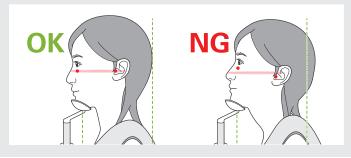
11. Check exposure position

The horizontal (A), front-back (B), and left-right (C) beams will intersect at the center of the exposure area.

Check that the beams are on target.



• If the line from the orbitale to the ear orifice is not horizontal, the back of the head might stick out and make contact with the arm during an exposure or ready move. For a patient who has a large head, run a test with no X-ray emission mode to make sure the arm does not contact the patient.



Tell the patient not to move his head during X-ray emission (or while the melody is playing) and then leave the X-ray room.



• Warn the patient not to move during emission (while the melody is sounding). If the patient moves, he could be hit by the arm or the image might be ruined.



Min. Scan Move Function

(If this function is turned on.)

12. Touch Ready key

Make sure the patient has his hands on the patient handle. With the unit already in the ready state, press the Ready key again.

The arm will move to its Min. Scan Move position and then a two-toned beep will sound.

After the arm moves, re-check the patient's positioning and hand position on the patient handle and then leave the X-ray room.

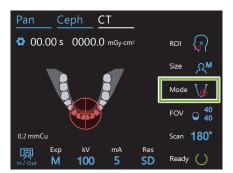
To continue go to "7.3.4 CT Exposure" (p. 81).

Relationship between CT and Two-Direction Scout Images

The CT and two-direction scout images are linked. If you double-click the green cross mark "+" in the two-direction scout image, the CT image for that place will be automatically displayed.

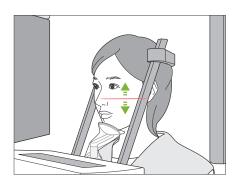
If the CT exposure has not been made, the cross mark "+" will be yellow. In this case, there is no linkage with anything.

7.3.3.3 Set FOV Position with Beam Positioning



When beam positioning is selected as the FOV positioning: "Mode", the FOV circle appears on the control panel. This shows the horizontal plane for the dental arch and is the approximate location of the exposure area. The front-back and left-right beams are linked to the FOV circle, and it will move accordingly when the beams are moved.

• The FOV circle determines a reference range, but its accuracy cannot be guaranteed.



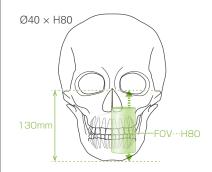
1. Line up horizontal beam

Move the horizontal beam up or down to set the vertical position. This beam indicates the height center of the FOV.

FOV Range

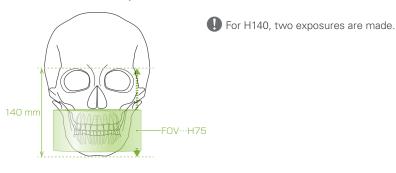
For Ø40 and Ø100:

The FOV has a vertical range of 130 mm. (It can be moved 50 mm up from its lowest position for an H80, 80 mm for an H50, and 90 mm an H40.)



For Ø150: * Ø150 only.

The FOV has a vertical range of 140 mm. (It can be moved 65 mm up from its lowest position for an H75, and 90 mm an H50.)



2. Line up front-back and left-right beams

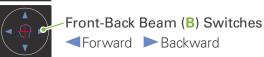
Ø40

The horizontal(A), front-back (B), and left-right (C) beams will intersect at the center of the FOV.

Move the front-back and horizontal beams to set FOV position.

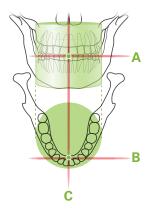
Check that the beams are on target.





● Ø80, Ø100, and Ø150

The horizontal (A), and left-right (C) beams will intersect at the center of the FOV. The front-back beam(B) is left lined up with the distal side of the left canine teeth and does not need to be adjusted. Check that the beams are on target.



* The FOV circle in the control panel is approximate. Check the beams for the exact position.

Tell the patient not to move his head during X-ray emission (or while the melody is playing) and then leave the X-ray room.

ACAUTION

• Warn the patient not to move during emission (while the melody is sounding). If the patient moves, he could be hit by the arm or the image might be ruined.

If the patient watches the arm go around, he may move his head. It is a good idea to tell the patient to close his eyes.

Min. Scan Move Function

(If this function is turned on.)

3. Press Ready switch

Make sure the patient has his hands on the patient handle. With the unit already in the ready state, press the Ready key again. The arm will move to its Min. Scan Move position and then a two-toned beep will sound.

After the arm moves, re-check the patient's positioning and hand position on the patient handle and then leave the X-ray room.

7.3.4 CT Exposure





1. Check ready state

Check that the Ready key on the control panel and the Ready LED (green) on the control box are on.



2. X-ray emission

Hold down the Emission switch.

The arm will rotate and X-ray emission will start.

The Emission LED (yellow) on the control box will light up and the melody will sound.

- * For the Ø150 × H140 exposure, two exposures are made one after the other. Continue to hold down the Emission switch after the first exposure is made. The arm will go up and return to its Ready position. Then the second exposure will be made. If you release the Emission switch before both exposures are completed, a complete image will not be made.
- ① As much as 15 seconds could elapse from the time the emission switch is pressed until X-ray emission actually starts. This is not abnormal. It takes some time for the unit to verify the computer settings.

MWARNING

• If an emergency occurs, release the Emission switch. Or press the Emergency Stop switch. This will stop the arm rotation and X-ray emission.

ACAUTION

- Go outside the X-ray room to press the Emission switch.
- If the operator must stay in the X-ray room for some reason, he should wear an X-ray protection apron and stay at least 2 meters away from the X-ray emission focal point. Also he should stay outside the area exposed to X-ray emission.
- Releasing the Emission switch during X-ray emission will stop the arm's rotation and terminate the exposure. In this case guide the patient away from the unit. If the X-ray head or the detector is directly in back of the patient, turn off the unit and slowly rotate the arm so the patient can leave. If you perform the ready procedures without having the patient move away from the unit, the arm could hit and possibly injure the patient.
- If an error occurs during an exposure, the arm will stop and the exposure will be terminated. In this case guide the patient away from the unit. If the X-ray head or the detector is directly in back of the patient, turn off the unit and slowly rotate the arm so the patient can leave. If you perform the ready procedures without having the patient move away from the unit, the arm could hit and possibly injure the patient.
- * If X-ray emission does not start when you press the Emission switch, check to see if the computer is busy reconstructing and image or asking for instructions.



3. Exposure completed

The melody stops when the exposure has been completed.

The arm will automatically move to the patient egress position and stop.

Release the Emission switch and hang it in its holder on the control box.

7.3.5 Patient Egress



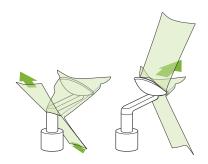
1. Guide patient away from the unit

Turn the temporal stabilizer knob and open the stabilizers completely.

Guide the patient away from the unit.

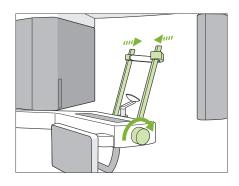
ACAUTION

- Take care that the temporal stabilizers do not strike the patient in the eye.
- Be careful of the stabilizers when the patient leaves the unit.



2. Throw away disposable covers etc.

Throw away the chinrest or bite-block cover. If the upper lip rest cover is removed by splitting its seams and turning it inside out, the outer surface will not touch anything.



3. Close temporal stabilizer

Turn the temporal stabilizer knob and close the stabilizers completely.



4. Touch Ready key

Touch the Ready key.

The "Arm return" dialogue will appear. Make sure the area is safe, and then touch the "Ok" button.

The arm will move back to its original patient positioning position.

* If another exposure will be made right away, touch the In/Out key to move the arm to its patient entry position.

7.3.6 Image Transmission



1. Image transmission

After the exposure is completed, the image is sent to i-Dixel. During transmission, the Ready LED on the control box is orange and blinks on and off.

After transmission has been completed, the Ready LED on the control box and the Ready key in the control panel will turn green and blink on and off.

① A two-toned beep sound when transmission has been completed and the LED turns green and blinks, but another exposure cannot be made until the image appears in the computer screen.





2. Image reconstruction

CT reconstruction will be performed.

- * Image reconstruction takes about 5 minutes.
- If all the CT tasks are save, the hard disk will fill up rather quickly. CT tasks should be moved or deleted periodically.
- ① During image acquisition, vertical or horizontal lines may appear. These are due to disparities in pixel sensitivity and does not indicate any sensor defect.
- Pror details on the software, refer to the user manual for i-Dixel.

7.4 Cautionary Remarks on Imaging

7.4.1 Artifacts Due to Sensitivity Discrepancies of Flat Panel Detector

The flat-panel detector (FPD) is an extremely dense and precise array of photo diodes (pixels). By compensating for the discrepancy in pixel sensitivity, the image can show greater detail than ordinary fluoroscopy. However, this discrepancy cannot be completely eliminated when a CT image is reconstructed.

To make a CT image, the X-ray beam rotates around the object and produces a fluoroscopic image. Points away from the center of the imaging area more on the imaging surface depending on the angle of their projection. This tends to even out the fluctuations in sensitivity, but it does not completely eliminate them. (See Fig. 1.) Therefore, artifacts can appear near the center of the imaging area as shown in Fig. 2. The appearance of these artifacts depends somewhat on the exposure conditions and the X-ray transparency of the object. To make a detailed analysis and precise diagnosis, the radiologist must carefully take into consideration the possibility and nature of artifacts.

Figure 1: Exposure Principle Diagram

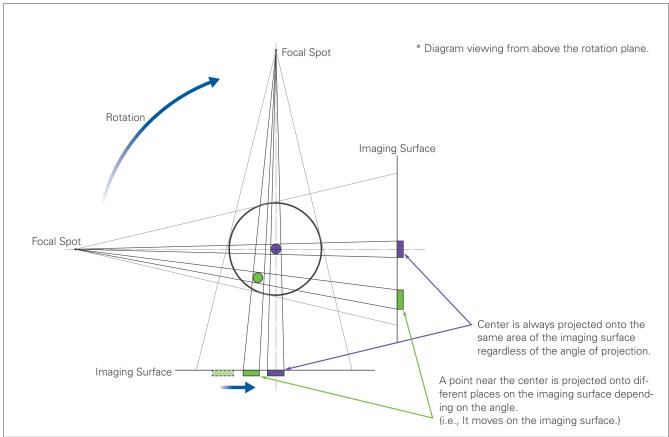
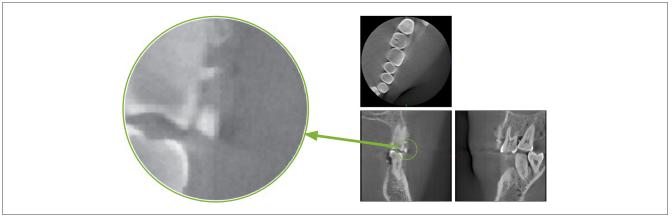


Figure 2: Example of an Artifact

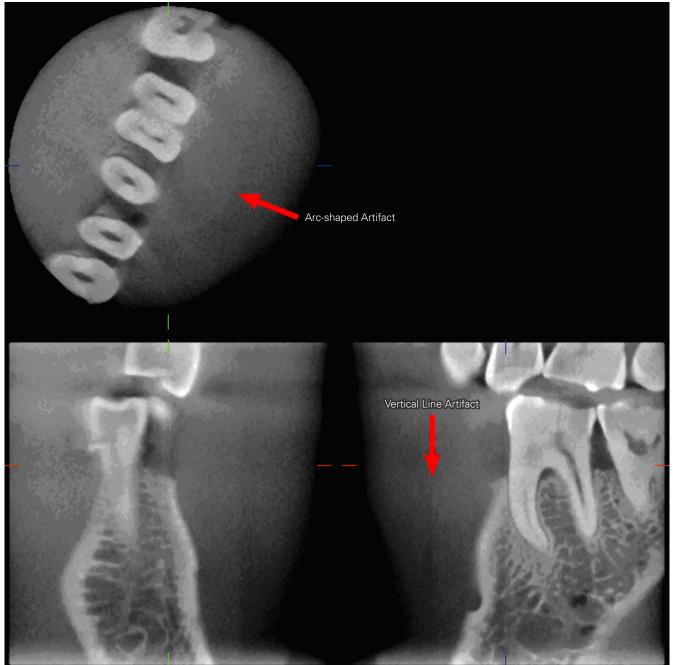


7.4.2 Artifacts Caused by Flat Panel Detector (FPD) Sensitivity Discrepancies

The flat-panel detector (FPD) is an extremely dense and precise array of photo diodes (pixels). Even though the panel is assembled in a clean room, even minutes deposits of dust or other debris can results in uneven discrepancies in the sensitivity of the photo-diodes. The image processing software compensates for these discrepancies and makes them all but invisible. However, logarithmic and filter processing during the reconstruction of the CT image may accentuate sensitivity discrepancies and produce artifacts in the CT image.

These discrepancies will have an arc shape in axial (z-plane) images and appear as vertical lines in coronal and sagittal images (x- and y-planes). (Fig. 3 below shows examples of these artifacts.) Artifacts can be identified by comparing the images of all 3 planes. If a vertical line in the coronal and sagittal planes is matched by an arc-shaped line in the axial plane, you may assume that these are artifacts.

Figure 3: Example of Artifacts



7.4.3 Artifacts Due to Angle of X-ray Penetration

Artifacts Characteristic of Cone Beam CT Imaging Artifacts depend on how far the object is from the plane of the X-ray beam's rotation.

CT images are a kind of tomography; images are reconstructed after collecting the images produced by the passage of X-rays emitted by a rotating beam. (Figure 4.1)

For cone beam CT imaging, artifacts appear above or below the plane of the beam's rotation because of insufficient information. These artifacts appear on a straight line running from the object to the source of the X-ray beam, and they are more conspicuous the farther the object is from the plane of the beam's rotation.

Figure 4: Track of X-ray Source and Detector

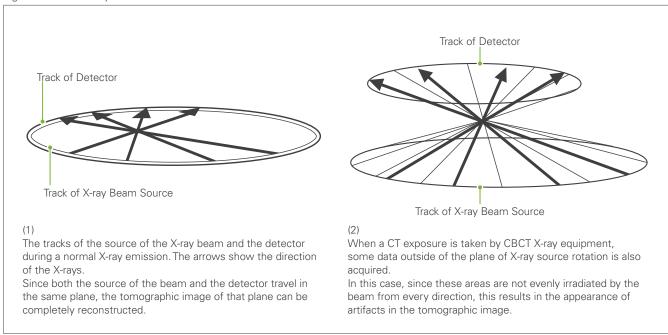


Figure 5 compares the case of 2 images of the same tooth made at different vertical positions. An artifact appears at the tip of the tooth because this area has a high CT value.

In Fig. 5.1, the artifact is nearly horizontal because it is at about the same height as the source of the X-ray beam.

In Fig. 5.2, the artifact is slanted because it is farther away from the plane of rotation for the beam source. It is also wider.

Radiologist must carefully take into consideration the nature of artifacts at the periphery of the area of interest.

Accurate Measurement of Thickness

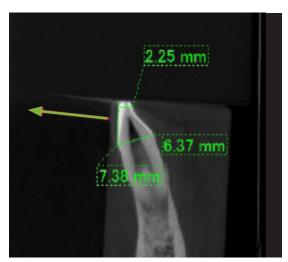
Following the principle stated above, the farther the area is from the plane of the beam's rotation, the less accurate the image is. This is demonstrated by measuring the thickness of a flat object positioned horizontally.

Figure 5.3 shows the images of aluminum disks with a thickness of 1.0 mm positioned 20 mm apart and parallel to the plane of the beam's rotation. All aluminum disks are arranged horizontally (parallel to the plane of X-ray source rotation). The bottom disk is 10 mm up from the bottom edge of the image. Therefore its thickness can be measured accurately. However, the measurement for the middle disk, which is 20 mm higher, is less accurate, and the measurement for the top disk is the least accurate of all.

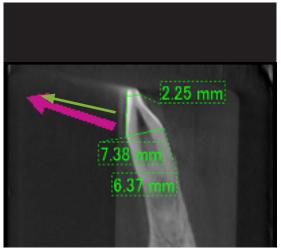
This phenomenon is occurred due to the artifacts, which are extending from each part of the aluminum plates overlapped, and these resulted making the boundary between the aluminum plates and air unclear.

This phenomenon must always be kept in mind for accurate diagnosis and analysis. Whenever an accurate measurement of thickness is required, adjust the bite block to position the area at the same height as the same height as the source of the X-ray beam.

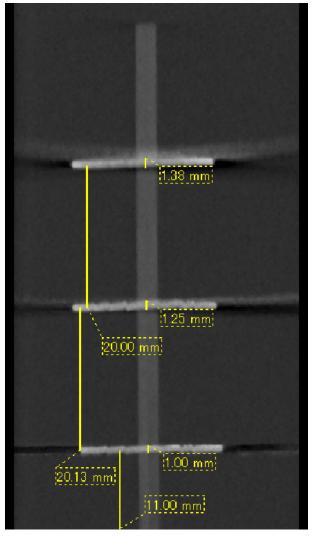
Figure 5: Artifacts



1. The artifact is produced near the plane of the beam's rotation



The area of the artifact is farther away, making it more conspicuous and giving it a different direction. Conspicuousness and direction of artifact are different.



3. CT image of three aluminum disks, each 1.0 mm thick, parallel to the plane of beam rotation but at different heights. The lowest one can be accurately measured because it is at the same height as the plane of the beam's rotation. The farther the disks are from the plane of the beam rotation, the thicker they appear in the image.

7.4.4 Artifacts due to Metal Prosthetics

Useful images may not be possible if a patient has metal fillings or prosthetic devices, and it is usually impossible to make a useful image of a crown if it is right next to a metal prosthetic.

Also it is sometimes impossible to make useful images of a root or jaw bone if there is a metal post, crown or other prosthetic right next to it.

Figure 6 through 12 were made of a mandible model with various metal prosthetics, posts and canal fillings to demonstrate what needs to be taken in consideration for accurate diagnosis and analysis.

Figure 6: Model of Full Metal Crown



Top of crown was removed and replaced with a piece of lead.

Figure 7: No Metal Prosthetic

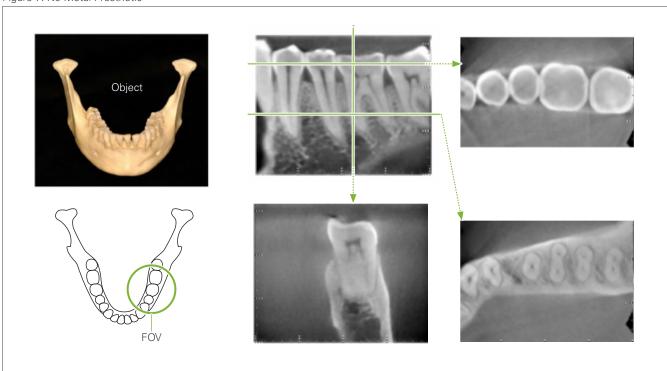


Figure 8: Full metal crown is on the opposite side of the FOV.

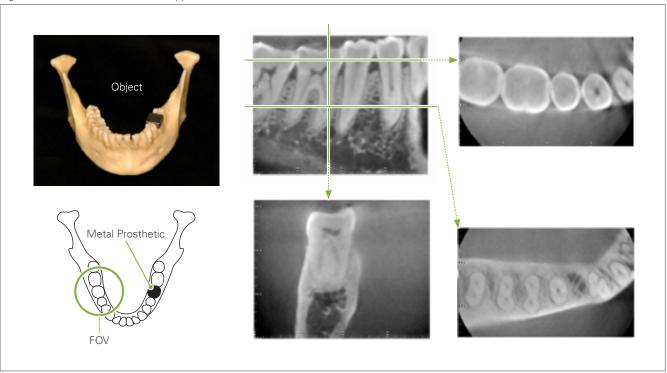


Figure 9: Full metal crown is on the same side of the FOV.

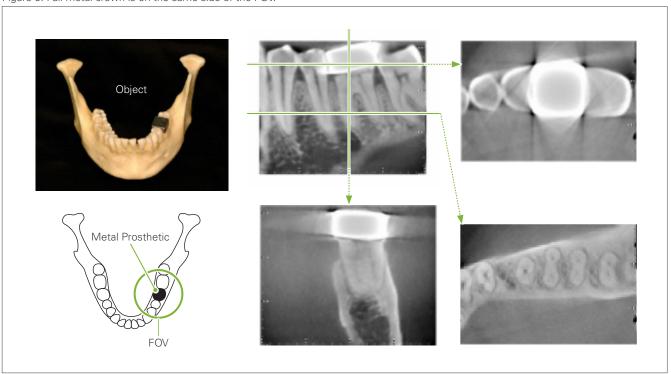


Figure 10 Post Crown



Left: Post and crown made after filling root canal with gutta-percha and point. Center: Post and crown attached to tooth.

Figure 11: Post crown is on the opposite side of the image area.

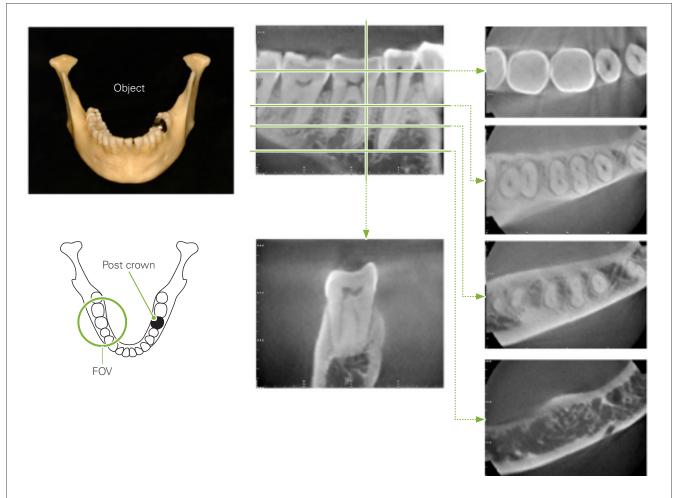
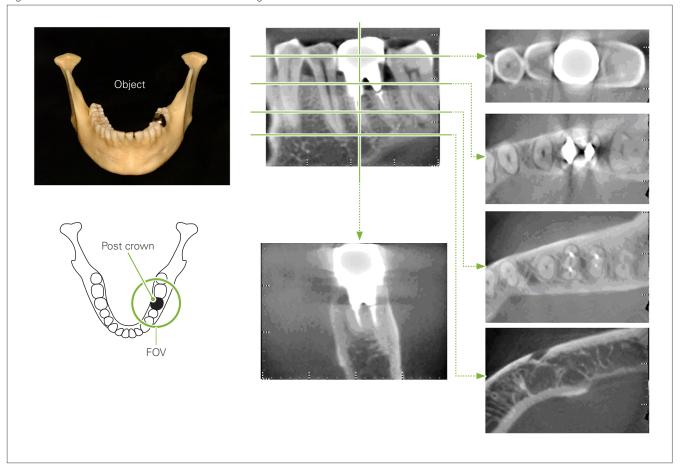
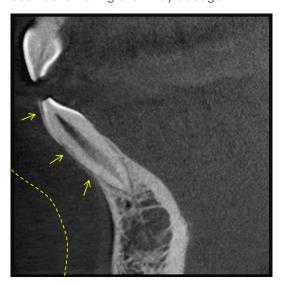


Figure 12: Post crown is on the same side of the image area.



7.4.5 Saturation Artifacts

If the X-ray beam is too strong, the flat panel sensor will be saturated, and this will affect the appearance of areas that absorb very little radiation such as soft tissues like lips, cheeks, and gums as well as thin hard tissues such as thin alveolar sockets or teeth. This must be kept in mind when using the image for diagnosis. This is especially a problem if there is a relatively large area of air; the dentist should consider steps such as lowering the X-ray dosage.



7.4.6 Artifacts for 180° Exposures

For 180° exposures, when the X-ray beam passes through the lower plane (a) in Figure 13, the result is the flat shape shown in Figure 14, where the beginning of 180° circuit matches the end of the circuit.

However, when the X-ray beam passes through the upper plane (b) in Figure 13, the result is the conical shape shown in Figure 15, where there is a discontinuity between the beginning and end of the circuit.

Therefore, some slight streaking appears in a 180° exposure that does not appear in a 360° exposure (Figure 16). These streaks run in the direction of the rotation. As a result, some semi-circular artifacts appear in the Z-plane image, and these must be recognized and taken into consideration for accurate diagnosis and analysis.

Figure 13

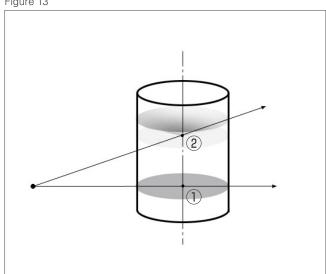
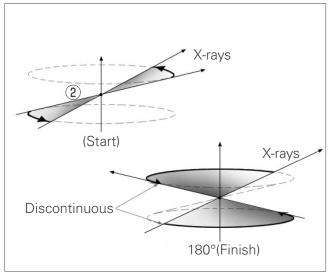
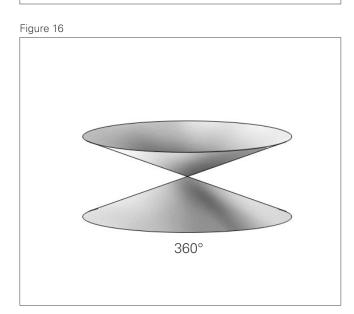


Figure 14 360° (Start) Continuous X-rays 180° (Finish)

Figure 15





7.4.7 Cautionary Remarks for Ø150 Exposures

Note the following cautionary remarks for Ø150 exposures, and take them into consideration when viewing the images.

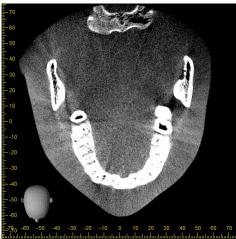
* For clarity of explanations, the artifacts shown in the images below have been highlighted by adjusting the contrast.

■ Ø150 Exposures

Streak Artifacts

In the axial plane, there may be artifacts that radiate out from the center of the FOV. (Figure 17)

Figure 17



Artifacts due to Density Disparities

For the X plane (back and forth) and the Y plane (left and right), there may artifacts due to a slight discrepancy in density. (Figure 18)

Figure 18





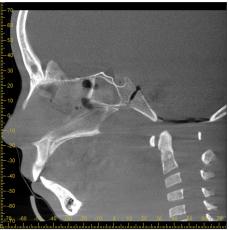
(a) X Plane (b) Y Plane

■ Ø150 × H140 Exposures

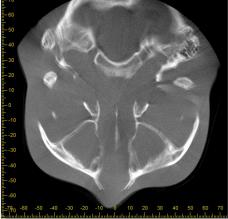
- Due to 2 different exposure data (top and bottom) are used, density might be uneven in the region where are joined.
- The accuracy of the shape of the X-ray target in the region where the 2 exposures are joined is not as good as a conventional CT scan. Therefore, the accuracy of measurements made in this region is also less accurate.
- If the patient moves between the top and bottom exposures, the images cannot be joined properly. (Figures 19 and 20) If the images are not properly joined, the following message will appear in i-Dixel. "Unable to determine the optimum stitching parameters.\r\nDefault values will be used instead. Check the resulting volume carefully when completed."

However, if the patient moves a lot, the message may not appear even though the images are not properly joined.

Figure 19: Poor Joining of Images



(a) X Plane



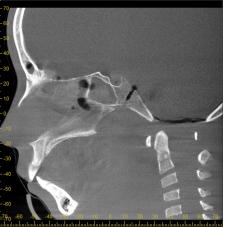
(b) Z Plane

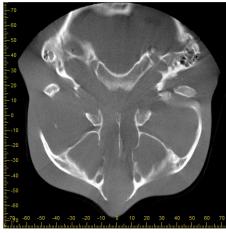
(a) A jag appears at the end of the nose

(b) Double exposure effect

If this happens only at the plane
where the images are joined in the
Z direction, the patient probably
moved in between the 1st and 2nd
exposures. If it happens throughout
the volume data, the patient probably moved frequently or constantly
during the exposure.

Figure 20: Normal Image





(a) X Plane (b) Z Plane

Corrections for Joint Position and Angle

If the patient moves between the 1st and 2nd exposures, the images will not match up properly when they are joined. Therefore, the position and angle will be automatically corrected based on the overlapped area of the original image. The degree of possible correction is noted below.

- The X, Y, and Z axes can be moved linearly up to 5 mm.
- The angles for the X, Y, and Z axes can be moved up to 3°.
- * These limits may be changed in the future.

Even within the limits noted above, it still may not be possible to join the images properly depending on how they are lined up. Also, it may not be possible to join the images depending on the X-ray target. Furthermore, if the images cannot be joined using the original images, they will be joined by using only the height information for the exposure.

• In order to correct for patient movement in between the 1st and 2nd exposures, the position and angle of the top exposure is automatically adjusted. For this reason a part of the top exposure may be missing as shown in figure 21 below. Figure 22 below shows an example of an image with a missing part.

Figure 21: Why a part is missing.

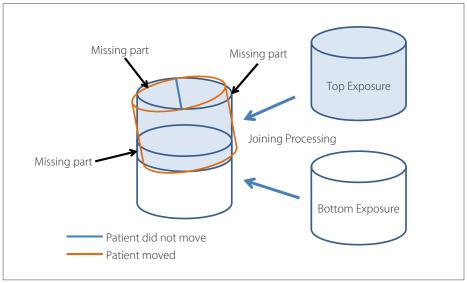


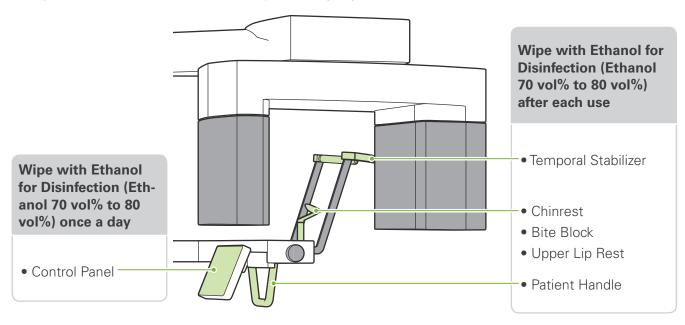
Figure 22: Example of Image with Missing Part



8 Maintenance, Parts Replacement, and Storage

8.1 Maintenance

Always turn off the Power switch before performing regular maintenance.



MWARNING

- Do not fail to turn off the Power switch. This will avoid the risk of electric shocks, burns, or accidentally pressing a switch.
- When disinfecting with Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%), take care that none of it seeps inside the unit. This could damage mechanical parts.
- Never use alkaline or acidic solutions, cresol soap, or other chemical solutions to clean outer surfaces. These can cause discoloration and material degradation.

 Use only Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) or a neutral detergent.
- Use Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%) to immediately wipe off any water, detergent or other chemicals that get on the outer surfaces.
- 1 The temporal stabilizers, chinrest, bite-block, upper lip rest, and patient handle cannot be autoclaved.

If it is not possible to obtain Ethanol for Disinfection (Ethanol 70 vol% to 80 vol%), use one of the disinfectants listed below; do not use any other type of disinfectant.

- DÜRR DENTAL's FD 322 quick disinfectant
- DÜRR DENTAL's FD 333 quick disinfectant
- DÜRR DENTAL's FD 360 imitation leather cleaning and care
- DÜRR DENTAL's FD 366 sensitive Rapid disinfection

8.2 Replacement Parts

- * Replace the parts as necessary depending on degree of wear and length of use. Refer to "12 Service Life, Consumables, and Replacement Parts" (p.111).
- * Order parts from local dealer or J. MORITA OFFICE.

8.3 Storage

Store the Veraview X800 under the following conditions:

Temperature: -10° C to $+50^{\circ}$ C ($+14^{\circ}$ F to $+122^{\circ}$ F); Humidity: 20% to 80% (without condensation); Atmospheric Pressure: 50 kPa to 106 kPa

- Do not expose the equipment to direct sunlight for an extended period of time.
- Store mouthpieces, disposable covers for the chinrest and bite-block, chinrest, and upper lip rest in a clean, hygienic place.
- If the unit has not been used for a long time, make sure it works properly before using.

9 Troubleshooting

9.1 Troubleshooting

If the equipment does not seem to be working properly, the user should first inspect the locations described below.

- * If the user is unable to inspect the equipment himself or if the equipment fails to work properly after being adjusted or after parts are replaced, contact your local dealer or J. MORITA OFFICE.
- * The inside parts of the equipment are charged with high voltage. Do not attempt to perform maintenance or adjustment that is not described in the troubleshooting table.

* If an accident occurs, the equipment must not be used until repairs have been completed by a qualified and trained technician provided by the manufacturer.

III MORITA

Before Inspection and Adjustment

Check that the Power LED (blue) on the control box is lit.

9.1.1 Main Unit

Symptoms	Possible Cause	Remedies
No power when the Power switch is turned on.	The Emergency Stop switch has been pressed.	Release the Emergency Stop switch and turn on the Power switch.
 Switches don't work. Control panel display doesn't respond. 	May still be responding to another key, or some process is in progress.	If normal operation is not restored after waiting a while, turn off the power, wait 1 minute and then turn it on again.
• The arm doesn't move.	Set in the Min. Scan Move.	Touch the Ready key to return it to normal ready state.
During usage: • Responds the same as when the Power switch is turned on. • Keys don't work.	NoiseMomentary power interruption	Turn off the Power switch and have the patient leave the unit. Turn on the Power switch again and see if it operates normally. * Make sure the power supply is on a devoted circuit and is AC 100 V, 20 A, minimum. Also the ground line must be sufficiently grounded.
 LCD display goes out. Display characters are corrupted. Display color is abnormal (blue etc.) Responds the same as when the Power switch is turned on. Keys don't work. 	Build up of static electricity.	Turn off the Power switch and have the patient leave the unit. Turn on the Power switch again and see if it operates normally. Make sure ground is properly connected. Maintain room temperature.

9.1.2 Exposures

Symptoms	Possible Cause	Remedies
Panorama & Cephalo Image is too light. Lines in image. White margins are too large. Image is partial. Image is completely black.	NoiseMomentary power interruption	Turn off the Power switch and have the patient leave the unit. Turn on the Power switch again and see if it operates normally. * Make sure the power supply is on a devoted circuit and is AC 100 V, 20 A, minimum. Also the ground line must be sufficiently grounded. Do not use any nearby devices that could produce noise during the exposure.
Panorama: Image has strobe effect.	Metal prosthetics can cause excessive feedback in the Auto Exposure system and produce a strobe effect. (AE Strobe)	Check by making exposure with the "Exp" set for "M" (manual).
Dental Arch Panorama: Incisor area is blurred.		Review "Positioning Point" guide.
Dental Arch Panorama: Center of image is white and left side is reduced.	Poor patient positioning.	Check that the front-back beam is lined up with the distal side of the left canine teeth.
Uneven density in image.	i-Dixel setting is not right.	Go to the i-Dixel tool bar and adjust this.
Futropolis dod,	Incorrect exposure conditions.	Auto Exposure: Adjust the auto level "Lv".
Extremely dark areas or the whole image is too white.		Manual Exposure: Adjust the tube voltage "kV" and current "mA".

9.1.3 i-Dixel

Symptoms	Possible Cause	Remedies
If the following error message appears in i-Dixel "Video capture failure."	LAN cable is not properly connected.	Make sure the LAN cable is properly connected and restart the computer.
If either of the following error messages appear in i-Dixel. "Cannot secure sufficient memory. Restart the software and the X-ray unit." "Cannot continue processing due to insufficient memory. Restart the software."	Insufficient computer memory due to repeated CT exposures or reconstructions.	Restart i-Dixel.
If the following error message appears in i-Dixel "Exposure terminated because no patient has been selected."	Exposure initiated without displaying a patient's image list.	Close the error message and then select and display a patient image list. Touch the Ready key to light it up. Then repeat the exposure procedure again.
If the following error message appears in i-Dixel "Exposure terminated because application is busy."	Initiated exposure when i-Dixel was not capable of processing a new exposure.	Close the error message and then select and display a patient image list. Touch the Ready key to light it up. Then repeat the exposure procedure again.

9.2 Error Messages

Error messages will appear in the control panel when some problem or error is detected. If the suggested response does not work, contact your local dealer or J. MORITA OFFICE. Make a note of the error number and report it when requesting help for the company.

Error No.	Messages
C1	The capture computer is not ready.
C2	The communication speed between the device and the HUB is insufficient.
C3	No response from the HUB.
C4	No response from the PC.
C5	The HUB between the device and capture system does not support jumbo frames.
C6	No response from the application software.
C7	The network interface card in the computer does not support jumbo frames.
01	The X-ray generator has overheated and acquisition has been stopped. Leave the power on and wait for approximately 30 minutes for the unit to cool down then try again.
02	Abnormal feedback detected in the voltage line of the X-ray generator. Acquisition has been stopped. Press OK to continue using the apparatus. If this message appears frequently, contact your local Morita representative for assistance.
03	A fault was detected in the fan motor of the inverter. Contact your local Morita representative for assistance.
04	Abnormal feedback was detected in the preheat line of the X-ray generator. Acquisition has been stopped. Press OK to continue using the apparatus. If this message appears frequently, contact your local Morita representative for assistance.
05	The tube voltage measured in the X-ray generator circuit was less than the specified value. Acquisition has been stopped. Press OK to continue using the apparatus. If this message appears frequently, contact your local Morita representative for assistance.
06	The tube voltage measured in the X-ray generator circuit was greater than the specified value. Acquisition has been stopped. Press OK to continue using the apparatus. If this message appears frequently, contact your local Morita representative for assistance.
07	The tube current measured in the X-ray generator circuit was less than the specified value. Acquisition has been stopped. Press OK to continue using the apparatus. If this message appears frequently, contact your local Morita representative for assistance.
08	An interlock error occurred in the X-ray generator. Acquisition has been stopped. Press OK to continue using the apparatus. If this message appears frequently, contact your local Morita representative for assistance.

Error No.	Messages
09	The overvoltage circuit has been activated. Acquisition has been stopped. Press OK to continue using the apparatus. If this message appears frequently, contact your local Morita representative for assistance.
0A	The overcurrent circuit has been activated. Acquisition has been stopped. Press OK to continue using the apparatus. If this message appears frequently, contact your local Morita representative for assistance.
0B	A PFC error occurred in the X-ray generation circuit. Acquisition has been stopped. Press OK to continue using the apparatus. If this message appears frequently, contact your local Morita representative for assistance.
0C	An abnormality was detected in the power supply control of the X-ray generator. Acquisition has been stopped. Press OK to continue using the apparatus. If this message appears frequently, contact your local Morita representative for assistance.
0D	The inverter circuit has overheated. Acquisition has been stopped. Leave the power on and wait for approximately 5 minutes for the unit to cool down then try again.
0E	Communication error between CPU and PCBs. The operation has been stopped.
OF	The arm and FPD control PCBs did not respond. Turn the unit off and on again. If this message appears frequently, contact your local Morita representative for assistance.
10	The X-axis and Y-axis motor control PCB did not respond. Turn the unit off then on again. If this message appears frequently, contact your local Morita representative for assistance.
11	The slit motor control PCB did not respond. Turn the unit off the on again. If this message appears frequently, contact your local Morita representative for assistance.
12	The operation panel control PCB did not respond. Turn the unit off then on again. If this message appears frequently, contact your local Morita representative for assistance.
13	The lift motor control PCB did not respond. Turn the unit off then on again. If this message appears frequently, contact your local Morita representative for assistance.
14	The inverter PCB did not respond. Turn the unit off then on again. If this message appears frequently, contact your local Morita representative for assistance.
15	The Cephalo cassette connector PCB did not respond. Turn the unit off then on again. If this message appears frequently, contact your local Morita representative for assistance.
16	The Cephalo control PCB did not respond. Turn the unit off then on again. If this message appears frequently, contact your local Morita representative for assistance.
17	Abnormality detected during arm rotation and movement has been stopped. Press the Ready button to return to the initial position. If this message appears frequently, the arm rotation mechanism or arm motor PCB may be faulty. Contact your local Morita representative for assistance.

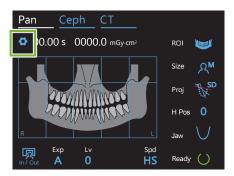
Error No.	Messages
18	Abnormality detected during X-axis operation and movement has been stopped. Press the Ready button to return to the initial position. If this message appears frequently, the X-axis motor or XY motor PCB may be faulty. Contact your local Morita representative for assistance.
19	Abnormality detected during X-axis operation and movement has been stopped. Press the Ready button to return to the initial position. If this message appears frequently, the Y-axis motor or XY motor PCB may be faulty. Contact your local Morita representative for assistance.
1A	Abnormality detected during operation panel support operation and movement has been stopped. Press the Ready button to return to the initial position. If this message appears frequently, the patient support elevating mechanism or frame motor PCB may be faulty. Contact your local Morita representative for assistance.
1B	Abnormality was detected in left/right beam operation and movement has been stopped. Press the Ready button to return to the initial position. If this message appears frequently, the left/right beam mechanism or frame motor PCB may be faulty. Contact your local Morita representative for assistance.
1C	Abnormality detected in horizontal slit left-side operation and movement has been stopped. Press the Ready button to return to the initial position. If this message appears frequently, the horizontal slit left side or slit motor PCB may be faulty. Contact your local Morita representative for assistance.
1D	Abnormality was detected in horizontal slit right-side operation and movement has been stopped. Press the Ready button to return to the initial position. If this message appears frequently the horizontal slit right side or slit motor PCB may be faulty. Contact your local Morita representative for assistance.
1E	Abnormality detected in vertical slit upper-side operation and movement has been stopped. Press the Ready button to return to the initial position. If this message appears frequently, the vertical slit upper side or slit motor PCB may be faulty. Contact your local Morita representative for assistance.
1F	Abnormality detected in vertical slit lower-side operation and movement has been stopped. Press the Ready button to return to the initial position. If this message appears frequently, the vertical slit lower side or slit motor PCB may be faulty. Contact your local Morita representative for assistance.
20	Abnormality detected during flat panel vertical movement. Press the Ready button to return to the initial position. If this message appears frequently, the arm rotation mechanism or arm motor PCB may be faulty. Contact your local Morita representative for assistance.
21	Abnormality detected during lift elevation and movement has been stopped. Press the up/down keys on the operation panel to try again. If this message appears frequently, the lift elevating mechanism or Z motor PCB may be faulty. Contact your local Morita representative for assistance.
22	Abnormality detected during X-ray generator rotation and movement has been stopped. Press the Ready button to return to the initial position. If this message appears frequently, the X-ray generator rotation mechanism or slit motor PCB may be faulty. Contact your local Morita representative for assistance.

Error No.	Messages
23	The second scout image was not captured.
24	Capturing the second scout image was interrupted.
25	Emission switch was released during image acquisition. Acquisition has been interrupted.
26	Emergency stop. Emission switch or a key on the operation panel was pressed during acquisition.
27	Returning the unit to its initial position was interrupted. Press the Ready button to return to the initial position.
28	Emergency stop. If a patient is positioned in the unit, have them exit the unit before proceeding. After confirming the unit may be operated safely, release the emergency switch by turning it to the right and pulling down.
29	Image data is not being received by the capture PC. Acquisition has been stopped. If this message appears at the start of the acquisition, this may indicate a timing issue. Please retry and if the error persists, contact your local Morita representative for assistance.
2A	Abnormality detected in Cephalo slit operation and movement has been stopped. Press the Ready button to return to the initial position. If this message appears frequently, the Cephalo slit or Cephalo PCB may be faulty. Contact your local Morita representative for assistance.
2B	Abnormality detected in Cephalo cassette operation and movement has been stopped. Press the Ready button to return to the initial position. If this message appears frequently, the Cephalo cassette or Cephalo PCB may be faulty. Contact your local Morita representative for assistance.
2C	Abnormal communication with 3DXD on the capture PC. The driver may not be running or may otherwise be occupied. Check the software status on the PC then try again.
2D	Abnormal communication with DixelD on the capture PC. The driver may not be running or may otherwise be occupied. Check the software status on the PC then try again.
2E	Abnormal communication with the digital cassette. Turn the unit off then on again and retry the operation.
2F	Scout position has been changed. Please set the scout position again from the capture PC.
30	Abnormality detected in copper filter switching mechanism operation and movement has been stopped. Press the Ready button to return to the initial position. If this message appears frequently, the filter changeover mechanism or slit motor PCB may be faulty. Contact your local Morita representative for assistance.
31	The equipment was moved after acquiring the scout image. The position is no longer valid. Please reacquire the scout image and try again.
32	The short connector of the X-ray generator is not connected.
33	The position is to high for stitched image acquisition. Use the up/down key to lower the arm and try again.

Error No.	Messages
34	The position is to low for stitched image acquisition. Use the up/down key to raise the arm and try again.
35	Abnormality detected in the arm vertical position data. Use the up/down key to lower the arm to the lowest position and reset the position data.
36	Image acquisition cannot be performed as the previous image is still being transferred to the capture PC. When the operation is finished, retry the acquisition.
37	The direction of the Cephalo craniostat does not match the Cephalo acquisition mode. Adjust either the mode or craniostat position then try again.
38	The temple stabilizers are open. Ensure that the temple stabilizers are closed when the arm is rotating to avoid interference.
39	Abnormality detected in the arm vertical position data. Use the up/down key to raise the arm to the highest position and reset the position data.
ЗА	GUI parameters are incorrect. Turn the unit off then on again.
3B	X-ray is not irradiated. If this message appears many times consecutively, the X-ray generator may be faulty. Contact your local Morita representative for assistance.
3C	Abnormal maximum height detected. Preform lift teaching to resolve. Move the arm to its lowest limit and continue to hold the down key for at least 5 seconds. The lifting teaching screen will appear. Follow the directions in the service manual.
3D	Acquisition has stopped due to an abnormality detected in the backup timer. Contact your local Morita representative for assistance.
3E	An abnormality was detected in the inverter PCB cooling fan. You can continue using the equipment, but take radiographs at intervals of at least 20 minutes. If no interval is provided, the inverter circuit might be overheated. If this message appears frequently, the PCB or the fan may be faulty. Contact your local Morita representative for assistance.
3F	An abnormality was detected in the X-ray head cooling fan. You can continue using the equipment, but take radiographs at intervals of at least 20 minutes. If no interval is provided, the X-ray head might be overheated. If this message appears frequently, the X-ray head or the fan may be faulty. Contact your local Morita representative for assistance.
40	The door to the x-ray room may be open. Please close the door completely before acquisition.
41	Abnormal communication with DixelD on the capture PC. The driver may not be running or may otherwise be occupied. Check the software status on the PC then try again.
42	The capture computer is not ready.

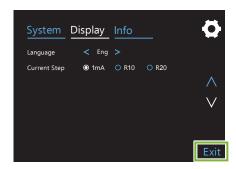
10 Various Settings

Enter Setting Display

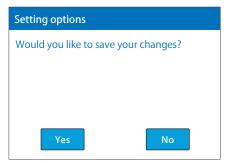


Touch the Settings key in the upper left corner. The display will change to the settings menu.

Exit Setting Display



Touch the Exit key in the lower right corner. The "Setting options" dialogue box will appear.

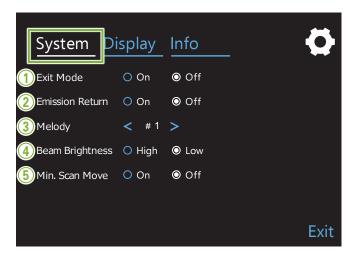


Yes: Touch this button to save the settings.

No: Touch this button to leave the settings as they are and return to the exposure screen. However, these settings will not be saved. The settings will go back to the previous ones when the Power switch is turned off.

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10.1 System Settings



Touch the System key to display a menu for various system settings for the unit.

The settings marked with a white circle () are the ones that are currently selected.

1. Exit Mode

This mode is to safely move the arm and lift to the secure place without making contact with any other device.

On: The arm moves to its specified position when the In/Out key on the control panel is held down.

ACAUTION

 Always make sure that the area is clear before doing this operation.

2. Emission Return

Press the Emission Switch once to return the arm after the patient has left the unit.



• Always make sure that the area is clear before doing this operation.

3. Melody

Select the melody to be played during emission.

- #1: Electronic beeping
- #2: Für Elise (Ludwig van Beethoven)
- #3: Minuet (Johann Sebastian Bach)
- #4: Symphony No. 9 (Ludwig van Beethoven)
- #5: Three Marches Militaires (Franz Schubert)
- #6: Hana "Flower" (Rentaro Taki)
- #7: Turkish March (Wolfgang Amadeus Mozart)
- #0: No melody

(A warning sound is emitted by the control box.)

4. Beam Brightness

Set the brightness for the beams.

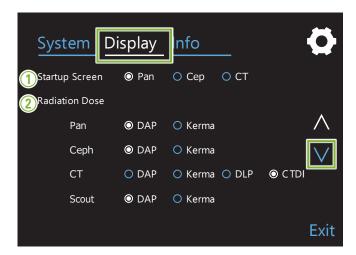
After making the setting, turn off the Power switch to activate it.

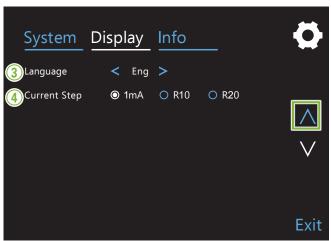
High: Bright Low: Dim

5. Min. Scan Move

After patient positioning, the arm can be moved to the position which will require the shortest exposure time (least time needed to hold down the Emission Switch) before the operator leaves the X-ray room.

10.2 Display Settings





Touch the Display key to switch the screen menu that shows possible settings for the display. Use the keys " \vee " or " \wedge " on the right side to change pages.

1. Startup Screen

Select the screen to be display when the unit is turned on.

2. Radiation Dose

Select the way to display the X-ray dosage.

3. Language

Select the language used for the control panel.

4. Current Step

This sets the increments for increasing or decreasing the "mA" for each time the key "+" or "-" is pressed.

R10 & R20 Settings

R10	R20
2.00	2.00
	2.24
2.50	2.50
	2.80
3.15	3.15
	3.55
4.00	4.00
	4.50
5.00	5.00
	5.60
6.30	6.30
	7.10
8.00	8.00
	9.00
10.00	10.00

10.3 Information

Touch the Info key to display information about the unit.

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11 Maintenance and Inspection

Regular Inspection

- * The user (i.e., clinic hospital etc.) is responsible for the management, maintenance and use of medical device.
- * Maintenance and inspection are generally consider to be the duty and obligation of the user, but if, for some reason, the user is unable to carry out these duties, he may rely on a qualified medical device serviceman. Contact your local dealer or J. MORITA OFFICE. for details.
- * This equipment should be inspected every 12 months in accordance with the following maintenance and inspection items. However, regular inspections may differs among countries, conduct the inspections accordance with the local regulation.
- * Replace the parts listed in the Regular Inspection List as necessary depending on degree of wear and length of use.
- * At the start and end of each business day, make sure that switching the Power switch to on and off turns the equipment on or off without fail.

	Category	Inspection Item
	FI 10 (.	Rated input voltage
	Electrical Safety	Ground connection
1		Power switch
'	Power or	X-ray emission and display
	Emergency Stop switches	Emergency Stop switch
		Arm emergency stop
		Floor and wall fastenings
2	Installation Integrity	Leveling
	2 Installation Integrity	Bolt and screw tightness
		All component installation
2	3 Positioning beams	Beam adjustment
3		Brightness
	4 Mechanical Operation	Arm rotation and moving parts
4		Lift function
		Wiring routing (cables)
5	Control Panel Display	Operation switches and keys
5	and Functions	Panel display
6	X-ray Head	Insulation oil leak
7	Outer Appearance	Patient positioning devices
	Outer Appearance	Outer surface
8	X-ray Control	Voltage control

	Category	Inspection Item
		Exposure field
		Uniformity
9	Panoramic Exposure	X-ray emission and image read-in
	Functions	Image
		Panorama left and right symmetry
		Exposure field
	Cephalometric Expo-	X-ray head direction change
10	sure Functions	Ear rod ring adjustment
	(option)	Uniformity
		Image
		Exposure Field
		Spatial resolution (IEC61223-2-6)
	CT Exposure	Noise gray scale and uniformity (IEC61223-2-6)
11	Functions and Performance	Contrast resolution (DIN6868-151)
		Image artifacts (DIN6868-151)
		Patient positioning (IEC61223-2-6)
		Basic functions
12	Computer Functions	Storage
		Network

12 Service Life, Consumables, and Replacement Parts

Service life refers to the standard period the unit or individual components can be expected to be usable as long as inspection and maintenance procedures specified by J. MORITA MFG. CORP. are followed.

Component Service Life List refers to components that can be expected to wear out, degrade or break depending on frequency and conditions of usage, which greatly affects how long these components retain their performance standards.

Consumables refers to parts and components that inevitably are degraded and need to be replaced periodically and are not covered by the warranty.

The product warranty is good for 3 years after delivery.

The components on the Component Service Life List that are noted "Yes" are critical for safety. These components must be inspected and replaced or have appropriate maintenance performed as necessary without fail before their standard service life expires.

The user must pay for parts and repairs that are performed after the warranty expires or the part has passed its specified standard service life. However, if a maintenance contract has been agreed to, this will depend on the contents of that contract.

For details concerning regular inspection and parts replacements, contact your local dealer or J. MORITA OFFICE.

Component Service Life List

Components	Standard Service Life	Critical Safety Component	Remarks
Moving Parts (for Arm and Lift)	45,000 exposures or 6 years whichever comes first.	Yes	Including cables, bearings, etc.
Motors (for Arm and Lift)	45,000 exposures or 6 years whichever comes first.	N/A	
X-ray Tube *1	15,000 exposures	N/A	
High Voltage Unit	3 years	N/A	
X-ray Detector (FPD) *2	3 years	N/A	
Printed Circuit Boards	6 years	Yes	
LCD Display	6 years	N/A	
Touch Panel, Operation Switches	3 years	N/A	
Patient Handle	6 years	Yes	
Temporal Stabilizer	3 years	N/A	
Ear Rod Plate	3 years	N/A	
Nasion Plate	3 years	N/A	

^{*1} The service life of the X-ray tube depends on the number and length of exposures it is used for as well as the output (tube voltage and current) and the time between exposures. Of these factors, the most critical is the number of exposures which degrades the anode. As the anode gradually degrades, stable output is lost, and the circuit protection system detects errors and terminates X-ray emission.

^{*2} The service life of the FPD mainly depends on the ambient conditions (temperature and humidity) where it is used and the accumulated amount of X radiation it receives. As the accumulated amount of received X radiation increases, the sensitivity of the detector is gradually degraded. High humidity can also lead to degradation. Semiconductor degradation caused by X radiation and the disparities for individual semiconductor units can cause parts of the detector to lose sensitivity. Loss of sensitivity can be remedied to some degree by performing sensitivity compensation and calibration during regular inspections, but partial sensitivity degradation cannot always be corrected.

Consumable Parts

Components	Code No.	Exchange Frequency	Critical Safety Component	Remarks
Mouthpiece	6270750	Single use	Yes	For infection control.
Bite-block Cover	6211120	Single use	Yes	For infection control.
Chinrest Disposable Cover	6215001	Single use	Yes	For infection control.
Chinrest	6215031	1 year or whenever scratched or damaged.	Yes	
Bite-block	6215041	1 year or whenever scratched or damaged.	Yes	
Upper Lip Rest	6215043	1 year or whenever scratched or damaged.	N/A	
Ear Rod	6290325	1 year or whenever scratched or damaged.	N/A	
Hand Plate	6293013	1 year or whenever scratched or damaged.	N/A	

Replacement Parts

Code No.	Description	Specification	Specification	Qu.
6112442	Main Fuse (AC 120 V unit)	25 A, 250 V	Fast-acting, High Breaking Capacity, Size 0.25 × 1.25 inches	1
6112473	Main Fuse (AC 220 – 240 V unit)	10 A, 250 V	Fast-acting, High Breaking Capacity, Size 5 × 20 mm	1

Fuse shall be replaced by qualified person. The user should never replace the fuse himself.

MWARNING

• Main fuse is on the Varistor Board located lower right side of the column where remains "live" even if the Power switch is turned off. Be sure to turn off the circuit breaker before servicing to avoid electrical shock.

Service

The Veraview X800 may be repaired and serviced by

- The technicians of J. MORITA's subsidiaries worldwide.
- Technicians employed by authorized J. MORITA dealers and specially trained by J. MORITA.
- Independent technicians specially trained and authorized by J. MORITA.

The circuit diagrams, component parts lists, descriptions, calibration instructions or other information will be available on request, only for the service personnel authorized by J. MORITA to repair those parts.

13 Technical description

13.1 Technical Specifications

Product Name	Veraview X800
Model	X800
Intended Use	Veraview X800 is intended to be used for panoramic tomography including linear tomography and scanogram, cephalometric radiography, and cone beam computed tomography. Veraview X800 is an extraoral source X-ray unit that is used for dental and head radiographic examination and diagnosis of teeth, jaw, oral structure, temporomandibular joint, skull including the dento-maxillofacial areas, and hand for maturity assessment, by exposing an X-ray image receptor to ionizing radiation. The device uses a cone shaped X-ray beam projected on to a flat panel detector, and the examined volume image is reconstructed to be viewed in 2D or 3D viewing stations. The device is to be operated and used by dentists and other legally qualified professionals for pediatric and adult patients.
Essential Performance	No X-ray irradiation without an active operation of the Emission switch. X-ray termination is done by releasing the Emission switch. No unexpected movement initiated without any active of operation.
Electric Shock Protection Classification	Class I
Electric Shock Protection Degree for Applied Part	Type B with applied component with component no conductive connection
Type B Applied Parts	Temporal Stabilizer, Chinrest, Bite-Block, Upper Lip Rest, Patient Handle Ear Rod, Nasion Plate, Hand Plate (for Cephalo unit)
Single Use Item	Mouthpieces, disposable covers for the chinrest and bite-block
Protection Against Ingress of Water	IPX0
Operation Mode	Non-continuous operation, momentary operation for U.S.A. and Canada
Operating Altitude	Max. 3,000 m
X-ray Head (including high voltage device)	
X-ray Tube	D-054SB or D-051SB
X-ray Tube Manufacturer	TOSHIBA or Canon
Nominal Focal Spot	0.5
Target Angle	5°
Focal Spot Material	Tungsten
Total Filtration	Inherent Filtration: Min. 2.5 mmAL, 75 kv/HVL 3.5 mmAL Added Filter: 0.2 mmCu (9.6 mmAL, 75 kV/HVL 3.5 mmAL) Indicated Location of Added Filter: Control Panel
Beam Quality(HVL)	Min. 3.6 mmAL at 100 kV Min. 3.2 mmAL at 90 kV Min. 2.9 mmAL at 80 kV Min. 2.5 mmAL at 70 kV Min. 2.2 mmAL at 60 kV

^{*}Specifications may be changed without notice due to improvements.

Ignition	Preheated		
Rectification	Direct current method		
Cooling Method	Oil cooling		
Leakage Radiation	1.0 mGy/h at 1 m or less		
Referrence Loading Conditions	Panoramic Exposures 90 kV, 540 mAs (90 kV, 9 mA with 1:59 duty cycle. e.g. 7.4 sec. emission with 7 min. 17 sec. intermission) CT Exposure 100 kV, 480 mAs (100 kV, 8 mA with 1:59 duty cycle. e.g. 9.4 sec. emission with 9 min. 15 sec. intermission) Cephalometric Exposure 100 kV, 600 mAs (100 kV, 10 mA with 1:59 duty cycle. e.g. 6 sec. emission with 5 min. 54 sec. intermission)		
Unit Specifications			
	60 – 100 kV in 5 kV increments Accuracy: Display value ±10% (Refer to each mode for details.)		
Tube Voltage	Netherlands	60 - 99 kV in 5 kV increments between 60 and 95 kV	
	Ukraine	60 - 90 kV in 5 kV increments	
Tube Current	2 – 10 mA in 1 mA, R10 or R20 increments Accuracy: Display value ±20% (Refer to each mode for details.)		
Exposure Time	Accuracy: Display value ±(5% + (Refer to each mode for details.)	50 ms)	
Reproducebility of Air Kerma	Coefficient of Variation: Max. 0.0	5	
Duty Cycle	1:59 (Maximum Irradiation Time: Time: 18 min. 12 sec.)	18.5 sec., Minimum Interval	
X-ray Field	Set automatically by select the ex	xposure mode.	
Dose Information	User selectable Pan, Cephalo, Scout: DAP (Dose Area Product) or Air Kerma CT: DAP (Dose Area Product), Air Kerma, CTDI (volume CT Dose Index), DLP (Dose Length Product) Displayed on control panel Accuracy: Display value ±50% * * Local criteria has priority if required in regulation.		
Primary Protective Shielding	2.2 mmPb (panorama, CT), 3.0 m	mPb (cephalo unit)	
Attenuation Equivalent	Exterior Cover in front of Imaging	Sensor: Max. 1.2 mmAL	
Patient Positioning Method	Laser, Panoramic scout, Two-Dire (Refer to each mode for details.)	ction scout	
Positioning Laser	Class 1 (IEC 60825-1: 2014) Max. 0.39 mW, 655 or 670 nm		
Emission Switch	Dead-man type		

Panorama Main Unit with Cephalo Unit		Long Column: W1,400 \times D1,200 \times H2,325 mm Short Column: W1,400 \times D1,200 \times H2,185 mm		
		Long Column: W2,000 \times D1,200 \times H2,325 mm Short Column: W2,000 \times D1,200 \times H2,185 mm		
Control Box		W120 × D60 × H120 mm		
Installation Space a	nd Weight			
		Panorama	with Cephalo Unit	
Installation Space		1.32 m ²	2.6 m ²	
Weight		Long Column: Approx. 190 kg Short Column: Approx. 185 kg	Long Column: Approx. 225 kg Short Column: Approx. 220 kg	
Power Requirement	nts			
Rated Input Voltag	e*1	AC 120 V	AC 220 – 240 V	
Frequency		50 – 60 Hz single phase	50 – 60 Hz single phase	
Line Voltage Regul	ation	Max. 8%	Max. 8%	
Range of Line Volta (Including Line Volta		AC 108 – 132 V	AC 198 – 264 V	
Innut	Operation	2.0 kVA	2.0 kVA	
Input	Standby	0.2 kVA	0.2 kVA	
Power Supply Apparent Resistance		0.5 Ohm	0.5 Ohm	
Over-Current Release at Distribution Panel*2		20 A, Earth leakage current breaker	15 A, Earth leakage current breaker	
Connection to Supply Mains		Permanent connection		
Maximum Line Current (only for U.S.A.)		18.5 A		
Technique Factor f	or the Maximum	Panorama, CT Model: 90 kV, 9 m	nA	
Line Current (only	for U.S.A.)	with Cephalo Unit Model: 100 k	V, 10 mA	
 WARNING To avoid the risk of electric shock, this equipment must only be connected to a supply mains 			ith protective earth.	
Environmental Data	1			
Operating Condition	ons			
Ambient Temperature Range		+10°C to +35°C (+50°F to +95°F)		
Humidity		30% to 75% (without condensation)		
Atmospheric Pre	ssure Range	70 kPa to 106 kPa		
Transport and Stor	age Conditions			
Ambient Temper	ature Range	-10°C to +50°C (+14°F to +122°F)		
Humidity		20% to 80% (without condensation)		
Atmospheric Pre	ssure Range	50 kPa to 106 kPa		

^{*1} Representative countries are:
• AC 120 V: U.S.A. and Canada
• AC 220 – 240 V: EU countries

Dimensions

^{*2} Circuit breaker shall be applied on distribution panel. It is recommended to wire a breaker only for this unit.

Panoramic Exposures			
Tube Current and Voltage	2 – 10 mA: 60 – 80 kV in 5 kV increments 2 – 9 mA: 85 and 90 kV		
Exposure Time	Refer to value in the control panel. Max. 18.5 sec.		
	Setting Range	-4 - 6 (11 steps)	
	Tube Voltage Range	60 – 90 kV	
Auto Exposure	Tube Current Range	2 – 10 mA	
Auto Exposure	Reproducebility of Air Kerma	Coefficient of variation max. 0.05	
	Auto exposure can only be used	for panoramic exposure.	
Magnification	1.3 ×		
Focal Spot to Image Receptor Distance	600 mm		
Image Quality	Line pair resolution: Min. 2.5 LP/ Low contrast resolution: Min. diameter 2.0 mm at HS, Mi		
CT Exposures			
	Standard: 2 – 10 mA: 70, 75, and 80 kV 2 – 9 mA: 85 and 90 kV 2 – 8 mA: 95 and 100 kV		
Tube Current and Voltage	X-ray Dose Reduction (DR): 3 – 10 mA: 70, 75, and 80 kV 3 – 9 mA: 85 and 90 kV 3 – 8 mA: 95 and 100 kV		
Exposure Time	Refer to value in the control panel. Max. 17.9 sec.		
Nominal Electric Power	0.8 kW (100 kV, 8 mA, 0.5 sec.)		
Field of View (FOV)	Ø40 × H40 mm, Ø40 × H80 mm, Ø80 × H40 mm, Ø80 × H50 mm, Ø80 × H80 mm, Ø100 × H40 mm, Ø100 × H50 mm, Ø100 × H80 mm, Ø150 × H50 mm, Ø150 × H75 mm, Ø150 × H140 mm		
Focal Spot to Image Receptor Distance	600 mm		
Image Quality	MTF: Min. 2 LP/mm at SD, Min.	2.5 LP/mm at HR	
Cephalometric Exposures (option)			
Tube Voltage	60 – 100 kV in 5 kV increments		
Tube Current	2 – 10 mA: Except density compexposure.	ensation is used for a lateral	
Tube current	Programmed Current Variability (tion is used.	Control: LA, density compensa-	
Exposure Time	Refer to value in the control pane	el. Max. 6.1 sec	
Magnification	1.1 ×		
Focal Spot to Image Receptor Distance	1,650 mm		
Image Quality	Line Pair Resolution: Min. 2.5 LP/mm Low Contrast Resolution: Min. diameter 2.5 mm		

^{*} Since panoramic exposures are image-layer exposures, distances measured on a panoramic image are not equal to the actual distances. * For tube voltage regulation in specific countries, see "Tube Voltage" in the "Unit Specifications" section.

13.2 Requirements for Computers or Other Devices Connected to Computers

1. The Veraview X800 has been tested and found to comply with the limits for medical devices to the IEC 60601-1-2:2007 for electro magnetic compatibility.

These limits are designed to provide reasonable protection against harmful interference in a typical medical installation.

This equipment generates uses and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to other devices in the environment. However, there is no guarantee that interference will not occur in a particular installation.

If this equipment does cause harmful interference to other devices, which can be determined by turning the equipment 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 device.
- Increase the separation between the equipment.
- Connect the equipment into an outlet on a circuit different from that to which the other device(s) are connected.
- Consult the nearest J. MORITA OFFICE, its representative or its dealer for help.
- 2. The following equipment connected to the analog and digital interfaces must be certified according to the respective IEC standards (i.e. IEC 60950-1 for data processing equipment and IEC 60601-1 for medical equipment).

Everybody who connects additional equipment to the signal input part or signal output part configures a medical electrical system, and is therefore responsible that the system complies with the requirements of IEC 60601-1. If in doubt, consult the nearest J. MORITA OFFICE, its representative or its dealer for help.

- * Some of the following devices may cause some technical problems with the Veraview X800. Ask your nearest J. MORITA OFFICE for proper selection of equipment and connections.
 - The following devices may not be located in the X-ray protection area or the patient environment except the Hub. If the Hub is conformed with IEC60601-1 or IEC60950-1 and enclosure leakage current is conformed with IEC 60601-1. After installation, check that the IEC 60601-1 leakage current levels are not exceeded.
- * If the above is not satisfied the following devices has to be located over 1,5 meters apart from the Veraview X800.
- * The operator shall not touch the LAN connector when touching the patient.
- * The Veraview X800 must be connected to it's own separate power supply. Computers or any other external devices must NOT be connected to the same power supply as the The Veraview X800.

WARNING

- connect only items that have been specified as part of medical electric system or specified as being compatible with medical electric system.
- Do not use multiple portable socket outlet nor extension cord for the system power supply.
- * Computers or any other external devices must be connected in accordance with IEC 60601-1.
- * Devices that do not meet the IEC 60950 standard (minimum requirements) must not be connected to the system as they may pose a threat to operational safety.
- * Computers or any other external devices must be cleaned in accordance with the manufacturer's instructions.
- * Computers or any other external devices must be transport, storage and operate in accordance with the manufacturer's instructions.

Other system requirements

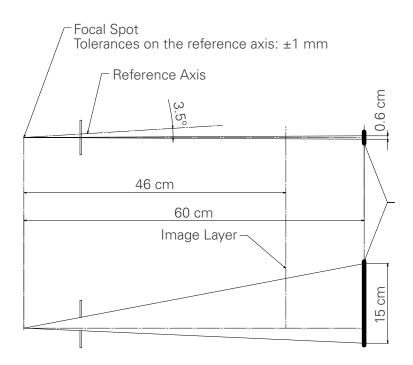
Hardware

CPU In Memory R. HDD H Video Board Re Network Protocol TC Network Interface U Others Nonitor 16	Alicrosoft Windows 7, 64 bit with Service Pack 2 Intel® Xeon or higher, or compatible IAM minimum 8 GB IDD 100 GB or more are recommended Idesolution of 1024 × 768 and color depth of 24bit ICP/IP with static IP address Iniversal 1 G BASE-T Ethernet NIC for internal connection Idetwork board, DVD drive Inch TFT LCD Identify Million colors Id		
Memory HDD H Video Board Re Network Protocol Network Interface Others N Monitor 17	AAM minimum 8 GB ADD 100 GB or more are recommended Albert 1024 × 768 and color depth of 24bit CCP/IP with static IP address Universal 1 G BASE-T Ethernet NIC for internal connection Altert 100 detection Altert 100		
HDD H Video Board Re Network Protocol TC Network Interface U Others N Monitor 16	IDD 100 GB or more are recommended desolution of 1024 × 768 and color depth of 24bit CP/IP with static IP address Universal 1 G BASE-T Ethernet NIC for internal connection detwork board, DVD drive 7 inch TFT LCD 6 Million colors 024 × 768 pixels or better EC60950-1 or IEC60601-1 EMC regulation delated UL standard (addition to U.S.A.)		
Video Board Network Protocol Network Interface Others Monitor Network Interface U 17	Resolution of 1024 × 768 and color depth of 24bit CCP/IP with static IP address Universal 1 G BASE-T Ethernet NIC for internal connection Retwork board, DVD drive 7 inch TFT LCD 6 Million colors 024 × 768 pixels or better EC60950-1 or IEC60601-1 EMC regulation Related UL standard (addition to U.S.A.)		
Network Protocol Network Interface Others Monitor To To To To To To To To To	CP/IP with static IP address Universal 1 G BASE-T Ethernet NIC for internal connection Jetwork board, DVD drive 7 inch TFT LCD 6 Million colors 024 x 768 pixels or better EC60950-1 or IEC60601-1 MC regulation Jelated UL standard (addition to U.S.A.)		
Network Interface Others N Monitor 17	Universal 1 G BASE-T Ethernet NIC for internal connection letwork board, DVD drive 7 inch TFT LCD 6 Million colors 024 x 768 pixels or better EC60950-1 or IEC60601-1 IMC regulation celated UL standard (addition to U.S.A.)		
Others N 17 Monitor 16	letwork board, DVD drive 7 inch TFT LCD 6 Million colors 024 x 768 pixels or better EC60950-1 or IEC60601-1 IMC regulation Related UL standard (addition to U.S.A.)		
Monitor 16	7 inch TFT LCD 6 Million colors 024 x 768 pixels or better EC60950-1 or IEC60601-1 IMC regulation Related UL standard (addition to U.S.A.)		
Monitor 16	6 Million colors 024 x 768 pixels or better EC60950-1 or IEC60601-1 IMC regulation Related UL standard (addition to U.S.A.)		
ır	MC regulation lelated UL standard (addition to U.S.A.)		
Standard EI Re	lelated C-UL standard (addition to Canada) ocal regulations		
Network Hub			
Specifications 5	000BASE-T compatible ports or more umbo Frame (Packet) supported		
Standard IE Re	EC60950-1 if it is used in non patient environment EC60601-1 or IEC60950-1 with leakage current conformed with IEC 60601-1. IMC regulation Related UL standard (addition to U.S.A.) Related C-UL standard (addition to Canada) EEE802.3x Local regulations		
Recommended for example N	IETGEAR GS105E		
Network Interface Card			
	ntel® PRO/1000 GT Desktop Adapter ntel® Gigabit CT Desktop Adapter		
Storage Devices D'	VD-R disk drive is recommended.		
Standard EI Re	EC60950-1 if it is used in non patient vicinity IMC regulation Related UL standard (addition to U.S.A.) Related C-UL standard (addition to Canada) Related regulations		
Other Equipment Connected to PC			
Standard EI Re	EC60950-1 if it is used in non patient environment IMC regulation Related UL standard (addition to U.S.A.) Related C-UL standard (addition to Canada) ocal regulations		
I AN Cania	Inshielded twisted pair cable with RJ-45 plug connections, over Cat. 5e, ength less than 2 m.		

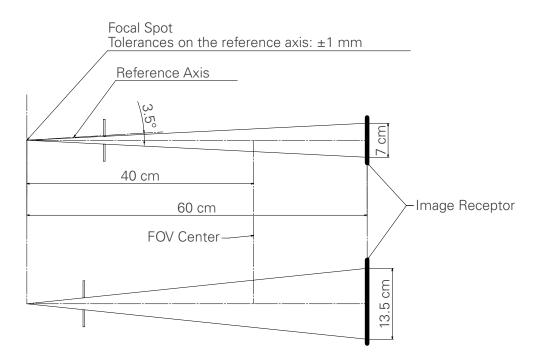
Application Software: Application software for image processing or database is provided by J. MORITA. MFG. CORP. It shall be used with Windows-based computer specified above. It conforms to 93/42/EEC (in EU), IEC62304 and 21 CFR (in U.S.A.), Medical device regulations (in Canada).

13.3 Relationship of Focal Spot, X-ray Beam and Image Receptor

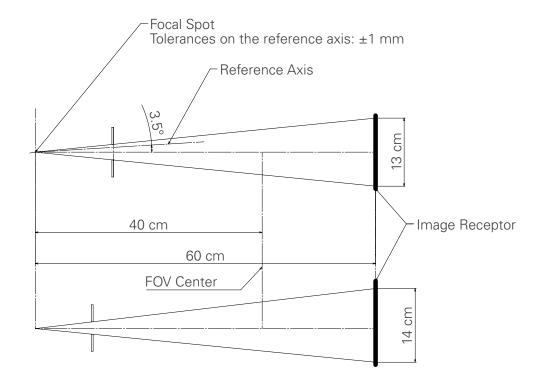
Panorama



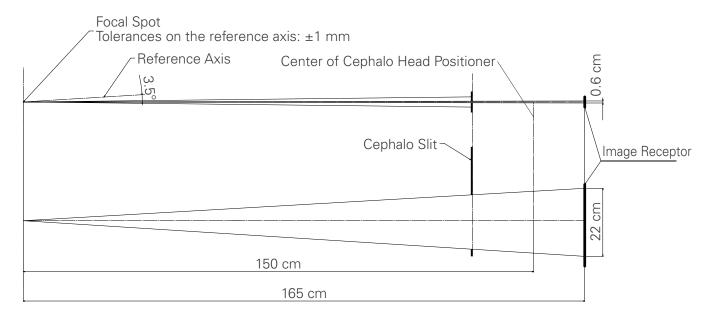
■ CT Ø40 × H80



CT Ø80 × H80



Cephalo

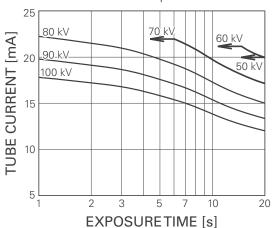


13.4 Tube Rating Chart

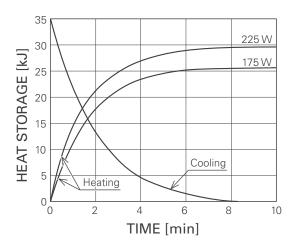
■ D-054SB

Maximum Rating Chart (absolute maximum rating chart)

Constant Potential High-Voltage Generator Nominal Focal Spot Value: 0.5



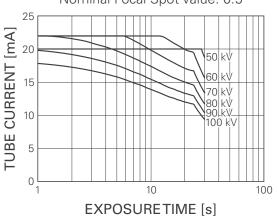
Anode Heating / Cooling Curve



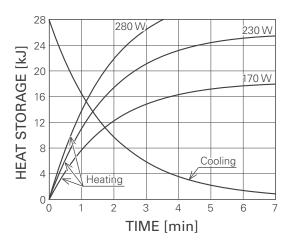
■ D-051SB

Maximum Rating Chart (absolute maximum rating chart)

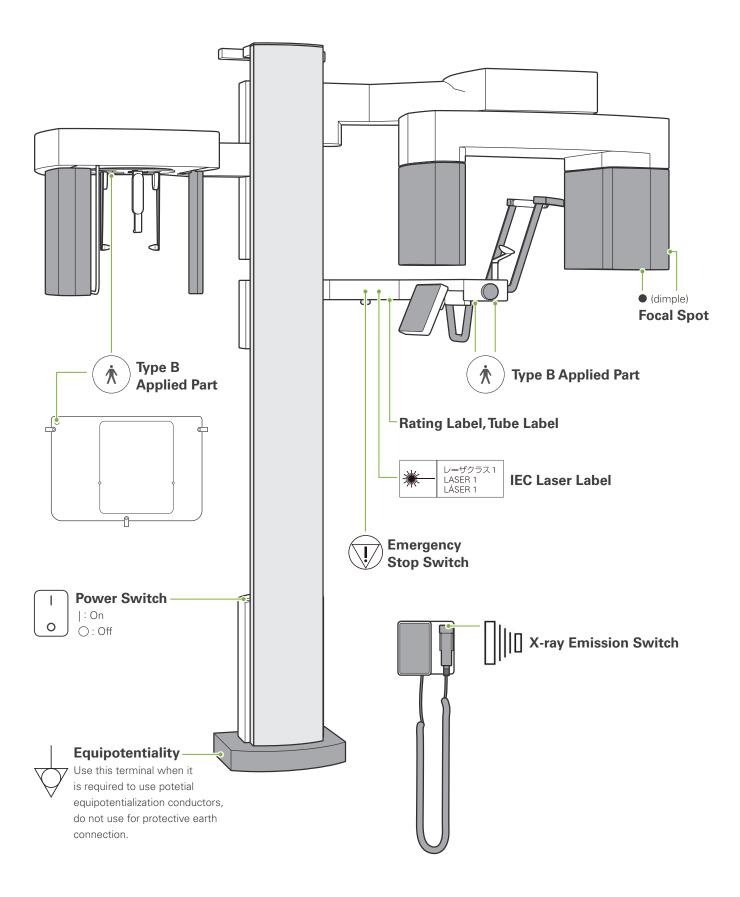
Constant Potential High-Voltage Generator Nominal Focal Spot Value: 0.5



Anode Heating / Cooling Curve



13.5 Symbols and Markings



■ Rating Label, Tube Label and Operation Instructions

* Some symbols may not be used.

Model: X800

Volt: 220 − 240 V ~

Input: 2 kVA

Stand By: 0.2 kVA Frequency: 50 – 60 Hz Duty Cycle: 1:59

Inherent Filtration: Min. 2.5 mmAL

Tube Voltage: 100 kV Tube Current: 10 mA

X-ray Tube Manufacture: TOSHIBA

X-ray Tube Model: D-054SB Nominal Focal Spot Value: 0.5

Head No.: Anode No.:

Registro Anvisa:

Registration number (Brasil)

Registro Sanitario:

Registration number (Mexico)

THIS EQUIPMENT COMPLIES WITH FDA RADIATION PERFORMANCE STANDARDS 21 CFR SUBCHAPTER J.: **Certification** (U.S.A.)



GS1 DataMatrix



Alternating Current



Date of manufactured



CE(0197) marking (EU)

Conforms with the European Directive,

0197 93/42/EEC.

CE marking (EU)

Conforms with the European Directive, 2011/65/EU.



WEEE marking (EU)

Conforms with the WEEE Directive



Attention, consult accompany documents.



cTUVus certification mark

(U.S.A. and Canada)



Manufacturer



Serial number



EU Authorized representative under the European Directive 93/42/EEC

(EU)

On Package



This way up

Fragile



Keep away from rain



Temperature limit



Humidity limit



Atmospheric pressure limit



Not Reusable

(Chinrest Disposable Cover etc.)

13.6 Standards and Procedures for Disposal of Medical Devices

Any medical devices which could possibly be contaminated must be first decontaminated by the responsible doctor or medical institution and then be disposed of by an agent licensed and qualified to handle standard industrial waste and industrial waste requiring special treatment.



This symbol indicates that the waste of electrical and electronic equipment must not be disposed as unsorted municipal waste and must be collected separately. Contact your local dealer or J. MORITA OFFICE. for details.

13.7 Description Based on Standards

21 CFR

• Caution: Federal law restricts this device to sale by or on the order of a dentist and a licensed healthcare practitioner. (for U.S.A.)

■ IEC 60601-1-3 and IEC 60601-2-63

- Test phantoms for dose information:
 Air condition (without any phantoms) for DAP, Air Kerma CTDI phantom, Diameter 160 mm in IEC 60601-2-44
- **Dosimeter:** Use the dosimeter calibrated without any phantoms according to the operation manuals of the dosimeter. Some dosimeter may calibrate with phantoms.
- Specified procedure: The jaw positioning tools (chinrest etc.) are removed and the temporal stabilizers are closed.
- Method used to estimate Air Kerma: A dosimeter (semi-conductor sensor) is put on the front outer cover of the image sensor at the center of the irradiation field. As for the distance conversion, the distance from the dosimeter to the surface of the image receptor must be considered (the distance from the outer cover to the image receptor is 12 mm). To measure it with an ionization chamber, the back scatter radiation must be subtracted.
- Method used to estimate dose area product: Measured by DAP (Dosage Area Product) meter. The DAP meter is attached to the front of the X-ray head for the Pan and CBCT, and in back of the secondary slit for the Cephalo. Be careful it does not fall and that its wiring is properly routed.
- Loading factor affecting to the patient dose: X-ray tube voltage, X-ray tube current, exposure time, X-ray field (FOV)
- Loading factor affecting to the radiation quality: Tube voltage
- Loading factor affecting to the value of reference air kerma: Tube voltage, Tube current
- Highest available air kerma condition: 100 kV, 10 mA
- Test instruction of X-ray tube voltage, current and exposure time: Constant (manual) exposure mode.
- Test instruction of Current time product: Multiple X-ray tube current and exposure time.
- Image processing applied to Original Data: Raw data send from the equipment is converted to original data for image reconstruction by compensation for dark image offset compensation, sensitivity compensation, saturation compensation, and defective pixel compensation.
- File transfer format of the images: The converted original data is saved as a "tif" file. Also, ancillary information such as dose etc. is save in text format. Reconstruction is performed based on the Original Data and appended information.

14 Appendix – Electromagnetic Declaration

This equipment conforms to IEC 60601-1-2: 2007, the relevant international standard for electromagnetic compatibility (EMC).

The following is the "Guidance and Manufacturer's Declaration" which is required by IEC 60601-1-2: 2007, the relevant international standard for electromagnetic compatibility.

MWARNING

- The Veraview X800 (hereafter referred to as the X800) needs special precautions regarding EMC and needs to be installed and put into service according to the EMC information provided in the ACCOMPANYING DOCUMENTS.
- Portable and mobile RF communications equipment can affect the X800.
- Use of parts other than those accompanied or specified by J.MORITA MFG.CORP may result in increased EMC emissions or decreased EMC immunity of the X800.
- The X800 should not be used adjacent to with other equipment. If adjacent use is necessary, the X800 should be observed to verify normal operation in the configuration in which it will be used.

Guidance and Manufacturer's Declaration - Electromagnetic Emissions

The X800 is intended for use in the electromagnetic environment specified below. The customer or the user of the X800 should assure that it is used in such an environment.

Emissions Test	Compliance	Electromagnetic Environment – Guidance	
RF emissions CISPR 11	Group 1	The X800 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		
Harmonic emissions IEC 61000-3-2	Class A *	The X800 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.	
Voltage fluctuations/flicker emissions IEC 61000-3-3	Clause 5 *	3	

^{*} This is only for AC 220 – 240 power supply.

Guidance and Manufacturer's Declaration - Electromagnetic Immunity

The X800 is intended for use in the electromagnetic environment specified below. The customer or the user of the X800 should assure that it is used in such an environment.

that it is used in such an environ	ment.		
Immunity Test	IEC 60601 Test Level	Compliance Level	Electromagnetic Environment – Guidance
Electrostatic discharge (ESD) IEC 61000-4-2	±6 kV contact ±8 kV air	±2, 4, 6 kV contact ±2, 4, 8 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 transients/bursts IEC 61000-4-4	±2 kV for power supply lines ±1 kV for input/output lines	±2 kV for power supply lines ±1 kV for input/output lines	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	±1 kV line(s) to line(s) ±2 kV line(s) to earth	±0.5, 1 kV line(s) to line(s) ±0.5, 1, 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 lines IEC 61000-4-11	<5% UT (>95% dip in UT) for 0.5 cycle 40% UT (60% dip in UT) for 5 cycle 70% UT (30 % dip in UT) for 25 cycle <5% UT (>95 % dip in UT) for 5 sec.	0% UT (>95% dip in UT) for 0.5 cycle 40% UT (60% dip in UT) for 5 cycle 70% UT (30% dip in UT) for 25 cycle 0% UT (>95% dip in UT) for 5 sec.	Mains power quality should be that of a typical commercial or hospital environment. If the user of the X800 requires continued operation during power mains interruptions, it is recommended that the X800 be powered from an uninterruptible power supply or a battery.
Supply frequency (50/60 Hz) magnetic field IEC 61000-4-8	3 A/m	3.15 A/m	Power frequency magnetic field should be at levels characteristic of a typical location in a typical commercial or hospital environment.
NOTE: UT is the a.c. mains v	oltage prior to application of th	e test level.	

Guidance and Manufacturer's Declaration - Electromagnetic Immunity

The X800 is intended for use in the electromagnetic environment specified below. The customer or the user of the X800 should assure that it is used in such an environment.

Immunity Test	IEC 60601 Test Level	Compliance Level	Electromagnetic Environment – Guidance
			Portable and mobile RF communications equipment should be used no closer to any part of the X800, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.
			Recommended Separation Distance:
	3 Vrms 150 kHz – 80 MHz	3.15 Vrms	$d = 1.2\sqrt{P}$
			$d = 1.0 \sqrt{P}$ 80 MHz – 800 MHz
	130 KHZ 00 WHZ		$d = 2.0 \sqrt{P}$ 800 MHz – 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).
D 11 1 1 DE	0.14		Field strengths from fixed RF transmitters, as determined by an electro-
Radiated RF	3 V/m	3.6 V/m	magnetic site survey, ^a should be less than the compliance level in each frequency range. ^b
IEC 61000-4-3 80 MHz – 2	80 MHz – 2.5 GHz		Interference may occur in the vicinity of equipment marked with the following symbol:

NOTE 1: At 80 MHz and 800 MHz, the higher frequency range applies.

NOTE 2: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

Recommended separation distances between portable and mobile RF communications equipment and the X800.

The X800 is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the X800 can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the X800 as recommended below, according to the maximum output power of the communications equipment.

Patad Maximum Output Payer	Separation Distance According To Frequency of Transmitter (m)			
Rated Maximum Output Power of Transmitter (W)	150 kHz – 80 MHz d = 1.2 \sqrt{P}	80 MHz – 800 MHz d = 1.0 √P	800 MHz – 2.5 GHz d = $2.0 \sqrt{P}$	
0.01	0.12	0.10	0.2	
0.1	0.38	0.32	0.63	
1	1.20	1.00	2.0	
10 3.79		3.16	6.32	
100	12.00	10.00	20.0	

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.

NOTE 1: At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.

NOTE 2: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

^a Field strengths from fixed transmitters, such as base stations for ratio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicated 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 X800 is used exceeds the applicable RF compliance level above, the X800 should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as reorienting of relocating the X800.

^b Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3.15 V/m.

Diagnostic and Imaging Equipment

Treatment Units

Handpieces and Instruments

Endodontic System

Laser Equipment

Laboratory Devices

Educational and Training Systems

Auxiliaries



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