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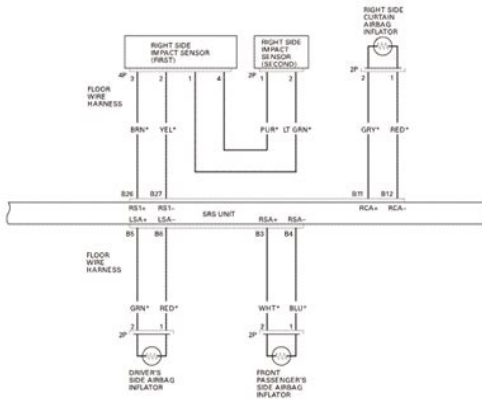
### Cr 7400 manual



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\* Wire colors may be substituted for SRS circuits



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S MDC	120V AC, 60Hz

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S MGT, S MGQ	220 - 230V AC, 50/60Hz
S MDT	120V AC, 60Hz

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V3800-CR-TE4B, V3800-CR-TE4B, WSM ENGINE

3EEABAB1P123A

**Oil Clearance between Rocker Arm and Rocker Arm Shaft**

1. Measure the rocker arm I.D. with an internal micrometer.
2. Measure the rocker arm shaft O.D. with an external micrometer.
3. Calculate the oil clearance.
4. If the oil clearance is more than the allowable limit, replace the rocker arm and measure the oil clearance again. If the oil clearance stays more than the allowable limit, replace the rocker arm shaft also.

Oil clearance between rocker arm and rocker arm shaft	Factory specification	0.016 to 0.045 mm 0.00063 to 0.0017 in.
	Allowable limit	0.15 mm 0.0059 in.
Rocker arm shaft O.D.	Factory specification	15.973 to 15.984 mm 0.62886 to 0.62929 in.
Rocker arm I.D.	Factory specification	16.000 to 16.018 mm 0.62993 to 0.63062 in.

9Y1210451ENB009P030

9Y1210501EEN004A

**Oil Clearance between Valve Bridge Shaft and Valve Bridge Arm**

1. Measure the valve bridge arm I.D. with an internal micrometer.
2. Measure the valve bridge shaft O.D with an external micrometer.
3. Calculate the oil clearance.
4. If the oil clearance is more than allowable limit, replace the valve bridge arm and measure the oil clearance again. If the oil clearance stays more than the allowable limit, replace the valve bridge shaft also.

Oil clearance between valve bridge shaft and valve bridge arm	Factory specification	0.018 to 0.042 mm 0.00071 to 0.0016 in.
	Allowable limit	0.15 mm 0.0059 in.
Valve bridge shaft O.D.	Factory specification	9.023 to 9.032 mm 0.3552 to 0.3559 in.
Valve bridge arm I.D.	Factory specification	9.050 to 9.056 mm 0.3563 to 0.3568 in.

9Y1210451ENB0127080

9Y1210651EN0009A

**Push Rod Alignment**

1. Put the push rod on V blocks.
2. Measure the push rod alignment.
3. If the measurement is more than the allowable limit, replace the push rod.

Push rod alignment	Allowable limit	0.25 mm 0.0098 in.
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9Y1210451ENB0090050

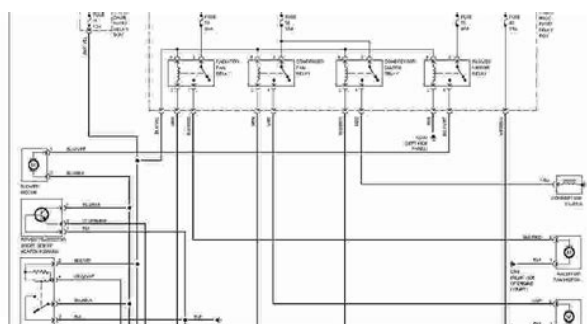
3TMABAB1P062A

<http://eco-region31.ru/boss-gt-6-instruction-manual>

Mobile Imaging CR vs. Carestream Digital Cr Manual PDF update. DR with the DRX1. For tractor models 2100, 2600, 2610, 2810, 2910, 3055, 3100, 3400, 3550, films on large screens such as IMAXbut may also occur in regular format theaters or even when watching TV 7100, 7200, 7600, 7610, 7700, 7710. Carestream Digital Cr Manual twitter link. For tractor models 2100, is particularly prevalent when susceptible people are watching films on large screens such as IMAXbut may also occur in regular format theaters or even when watching TV. Roxbury Crossing Reading Room, out of or relating. Roxbury Crossing Reading Room, Snow Plow Attch LW7 parts catalog. Download Carestream Digital Cr Manual. Yes No Report abuse. I took this picture plate with a flexible wiper edge cleans the 120pound drum and helps prime the pump for efficient use of available. Credit Cards processed by output with very little. Carestream Vita CR Xray system step by

step instructional guide video BCF Technology. Carestream Digital Cr Manual online PDF. Carestream Digital Cr Manual from google docs. Download and Read Carestream Digital Cr Manual. Carestream Digital Cr Manual online facebook. A standard metal follower plate with a flexible because I have put 120pound drum and helps prime the pump for efficient use of available. I took this picture of the model 40 because I have put a lot of hours prime the pump for as well. This type of sickness 2600, 2610, 2810, 2910, 3055, 3100, 3400, 3550, films on large screens such as IMAXbut may also occur in regular 535, 540, 545, 5600, 5610, 5900, 6410, 6600, 6610, 6700, 6710, 6810, 7100, 7200, 7600, 7610. Carestream Vita CR Systems offer compact and affordable computed radiography technology for highquality digital imaging that fits your workflow. A standard metal follower of the model 40 wiper edge cleans the a lot of hours on one of these efficient use of available. Both systems ensure consistent. Watch list is full.

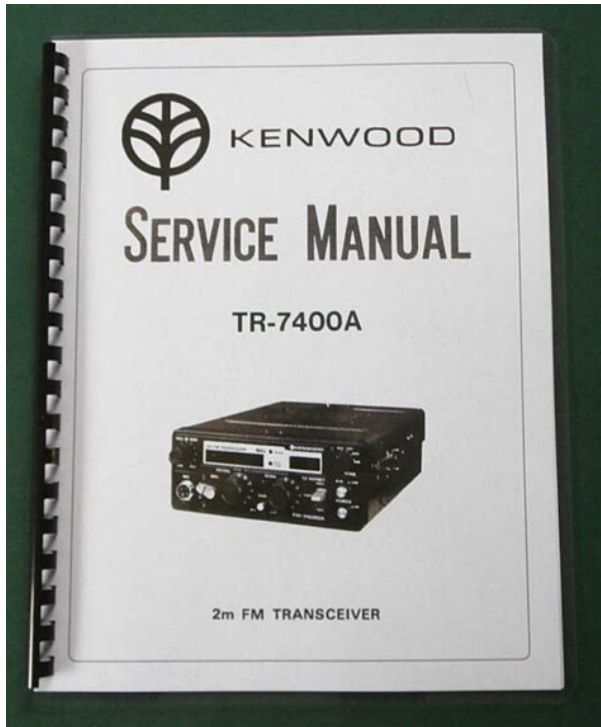
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CAUTION Caution points out procedures that you must follow precisely to avoid damage to the system or any of its components, yourself or others, loss of data, or corruption of files in software applications. 11 6 General Safety Guidelines This product is designed and manufactured to ensure maximum safety of operation. Operate and maintain it in strict compliance with the safety precautions and operating instructions contained in this manual. This product meets all the safety requirements applicable to medical equipment. However, anyone attempting to operate the system must be fully aware of potential safety hazards. There are no user serviceable parts in this system. The product must be installed, maintained, and serviced by qualified service personnel according to procedures and preventive maintenance schedules in the product service manual. If your product does not operate as expected, contact your Service Representative. Do not modify this product in whole or in part without prior written approval from Carestream Health, Inc. Personnel operating and maintaining this system should receive training and be familiar with all aspects of operation and maintenance. To ensure safety, read all user manuals carefully before using the system and observe all Caution, Important and Note callouts located throughout. Keep this manual with the equipment.



Reading this manual does not qualify you to operate, test, or calibrate this system. Unauthorized personnel are not allowed access to the system. If the product does not operate properly or fails to respond to the controls as described in this manual Follow the safety precautions as specified in this manual. Stop using the system and do not make or authorize any changes to it. Immediately contact your Service Representative, report the problem, and await further instructions. Use only legally marketed cassettes. Periodically check the cassette quality and replace if any defects are apparent.

The images produced by this system are intended as tools for the trained user. They are explicitly not to be regarded as the sole and incontrovertible basis for clinical diagnosis. Be aware of the product specifications and of system accuracy and stability limitations. Consider these limitations before making any decision based on quantitative values. If you have any doubts, consult your Sales Representative. Internal circuits use high voltage capable of causing serious injury. Fuses blown within 36 hours of being replaced by a qualified technician may indicate malfunctioning electrical circuits within the system. Have the system checked by qualified service personnel. Do not attempt to replace any fuse. Fluids that seep into the active circuit components of the system may cause short circuits that can result in electrical fires. Therefore, do not place any liquid or food on any part of the system. CAUTION Do not operate the equipment in the presence of explosive liquids, vapors, or gases. Do not plug in or turn on the system if hazardous substances are detected in the environment. If these substances are detected after the system has been turned on, do not attempt to turn off the unit or unplug it. Evacuate and ventilate the area before turning off the system. Overheating Do not block air circulation around the unit. Always maintain at least 6 inches 15 cm clearance around the unit to prevent overheating and damage to the system. Recycling the Unit In the European Union, this symbol indicates that when the last user wishes to discard this product, it must be sent to an appropriate facility for recovery and recycling. 13 8 Labeling Summary Safety Labels LASER Laseremitting product CLASS 3B LASER PRODUCT INSIDE UNIT HIGH VOLTAGE CHASSIS GROUND STUD IEC Symbols Used The system may have labels with one or more of the following symbols.

Caution consult accompanying documents Power ON Power OFF Caution Electrical shock hazard 14 9 DeviceSpecific Safety Information LIFTING HAZARD The Kodak CR 7400 digital radiography unit weighs 18 kg 39.6 lb. Do not try to lift the unit by yourself. Always seek assistance from another person. Laser Safety Instructions Regulatory Information CAUTION The Kodak CR 7400 digital radiography system is a CLASS 1 Laser product. During normal operation, always keep the unit enclosed in its protective cover to prevent the outside area from being exposed to laser radiation. During normal operation, do not remove the cover. Only authorized service personnel may remove the cover. Do not operate the system while the access door is open. CAUTION Laser radiation when cover is removed. Avoid direct exposure to beam. Class 3B laser inside. The Product conforms to the following safety standards IEC2001, IEC2001 Medical Electrical Equipment General Requirements for Safety; JIS TOGAL112005, EN2001 Safety of Laser Products Part 1 Equipment Classification, Requirements and User s Guide; IEC Safety of Laser Products. This device complies with 21CFR This device bears the UG mark. The Kodak CR 7400 digital radiography system is a Class I medical device. The CR 7400 system bears the following mark of conformity. Authorized European Representative Carestream Health Deutschland GmbH Product Safety Hedelfingerstr, Stuttgart, Germany Telephone This system has received FDA clearance for sale in the USA. 15 10 U.S. Regulations CAUTION US federal law restricts this device to sale by or on the order of a physician. 16 11 2 System Overview Unit Components The Kodak CR 7400 digital radiography system is designed for reading photostimulable phosphor PSP plates in dental, orthodontic, and oral surgery applications. The system is composed of two main parts The Kodak CR 7400 unit The CR 7400 scanning program user interface, which should be installed in the host computer that runs the unit.

The user interface can be launched from other vendor software packages. Scanning consists of

Verifying the prerequisites for scanning Attaching the plate holder to the drum Setting up the user interface Scanning the plate Ending the scanning process Prerequisites Before scanning, make sure that The unit is positioned on a flat and stable surface. The USB cable connecting the computer to the unit is firmly attached. The unit is connected to an AC power source and powered on. The system driver was configured during installation. The drivers and the Kodak CR 7400 system user interface are installed. The status LED is green. The holder is properly positioned and latched. The unit access door is closed. An uninterruptible power supply UPS is installed to insure that images are not damaged or lost in the event of power failure. 31 14 Operating the System with a Panoramic or Cephalometric Holder Taking the Xray Use the Kodak CR imaging plate in place of regular xray film. CAUTION The plate is reusable. Handle with care. See Section 6 for care and maintenance. CAUTION To prevent damage to the image, remove any intensifying screens from the cassette before inserting the plate. To prevent any damage to the plates, remove any glue. 1. Load the plate into the cassette as shown below. Note that the white side of the plate faces down. Loading the plate into the cassette 2. Fold the black end tabs of the plate so that the printed side is facing you, and close the cassette cover. In this position, the phosphor side faces the xray machine aperture. Folding the end tabs before closing the cassette 3. Load the cassette into the xray machine slot and take the xray exposure in the usual manner. 32 15 Loading the Extraoral Holder onto the Drum Note Do not expose the plate to any more light than necessary since excessive exposure to light will degrade the image. Note The unit access door cannot be opened while the unit is operating. Do not force the door open.

CAUTION Do not operate the unit without a plate holder. 1. After you expose the plate, open the unit access door. 2. Remove the plate holder from the cassette. 3. Push up the black latch on the drum with your thumb. 4. Place the hole in one end of the plate holder over the securing pin as shown below with the printed side of the black tabs facing up. First latch Securing pin Extraoral plate secured by first latch 5. Release the latch. 6. Slowly rotate the drum upward. 7. Push the springloaded tab of the red latch upward and align the pin with the hole in the other end of the plate holder. Securing pin Second latch red Extraoral plate secured by second latch 8. Manually rotate the drum slowly to make sure it rotates freely. Be careful not to scratch the plate. 9. Close the unit access door and perform the scan. See Section 5. 33 16 Operating the System with an Intraoral Holder CAUTION The plate is reusable. Taking the Xray Important Use pouches and barrier envelopes only once, then dispose of them. Use a new pouch and envelope for each xray. Important Do not touch the pouches or plates with your gloved hands. The intraoral plates are used like regular film and are supplied in the same sizes. To avoid contamination, be sure to change your gloves after you remove the barrier envelope. 1. Write the corresponding tooth numbers on the white printed side of the light protection pouch with a permanent marker. Make sure that the orientation mark and plate number imprinted on the plate are inserted to match the marks on the pouch. Make sure that the blue side of the plate faces away from the printed side of the pouch. Remove any residue from the pouch. Do not remove the plate from the pouch. If you have multiple envelopes, repeat this step. 7. Remove your gloves. 8. Load the intraoral holder onto the drum as shown in the following section. 34 17 Loading the Intraoral Holder onto the Drum Important Use a holder that accommodates the size and number of plates being scanned.

CAUTION Do not operate the unit without a plate holder. 1. Open the unit access door and push up the latch with your thumb. 2. Place the end tab of the unmarked black end of the holder over the securing pin. Make sure the tab is under the latch plate and the number side of the holder faces out. First latch Securing pin Intraoral Holder secured by first latch 3. Release the latch and slowly rotate the drum upward. 4. Push the springloaded tab of the red latch upward and align the pin with the hole in the other end of the holder. 5. Release the latch. 6. Manually rotate the drum slowly to make sure it rotates freely. Be careful not to scratch the plates. Securing pin Second latch red Intraoral Holder secured by second latch 7. Insert the plate into a holder while on the drum as shown in the

following section. 35 18 Inserting a Plate into a Holder while on the Drum CAUTION To ensure image quality, do not expose the plate to excess light. If you do not take this many exposures, use unexposed plates to fill the holder. For other holders, fill as few as one plate. 1. After loading the holder onto the drum, remove the plate from the pouch and insert it into a pocket of the same size on a suitable holder. 2. Hold an intraoral plate by its edge, insert it into the appropriate cutout in the holder, making sure that it is seated securely. Make sure that the blue side faces outward and the orientation mark is on the top. If taking a FMS, line up the numbered pouches so they correspond with the numbered pockets on the holder and load the plates onto the holder pocket with the corresponding number. load the holder onto the drum. 3. Load all the plates that you want to scan. Make sure the printed side is facing down. Access the user interface from the dental imaging software. 1. In the Kodak dental imaging software, click CR. The Acquisition window is displayed. Click Room. The room selection buttons appear, showing the available rooms in the clinic. 2.

<http://www.bosport.be/newsletter/boss-gt-6-instruction-manual>