

FOR OFFICE USE ONLY (DATE RECEIVED)	COLORADO DEPARTMENT OF PUBLIC HEALTH & ENVIRONMENT HMWMD - X-Ray Certification Program 4300 Cherry Creek Drive South, B1 DENVER, COLORADO 80246-1530 (303) 692-3448 or 3443 FAX (303) 691-7841 REPORT OF ASSEMBLY OF A DIAGNOSTIC X-RAY SYSTEM	CDPHE - FORM 2579
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1. EQUIPMENT LOCATION

2. ASSEMBLER INFORMATION

a. NAME OF HOSPITAL, DOCTOR OR OFFICE WHERE INSTALLED		a. COMPANY NAME	
COLORADO FACILITY REGISTRATION NO.:		COLORADO SERVICE COMPANY REGISTRATION NO.:	
b. STREET ADDRESS		b. STREET ADDRESS	
c. CITY	d. STATE	c. CITY	d. STATE
e. ZIPCODE	f. TELEPHONE NUMBER	e. ZIP CODE	f. TELEPHONE NUMBER

3. GENERAL INFORMATION

a. THIS REPORT IS FOR ASSEMBLY OF CERTIFIED COMPONENTS WHICH ARE (CHECK APPROPRIATE BOX(ES))		
<input type="checkbox"/> NEW ASSEMBLY - FULLY CERTIFIED SYSTEM	<input type="checkbox"/> NEW ASSEMBLY - MIXED/UNCERTIFIED	
<input type="checkbox"/> REASSEMBLY - FULLY CERTIFIED SYSTEM	<input type="checkbox"/> REPLACEMENT COMPONENTS IN AN EXISTING SYSTEM	
<input type="checkbox"/> REASSEMBLY - MIXED SYSTEM (Both certified and uncertified components)	<input type="checkbox"/> AN ADDITION TO AN EXISTING SYSTEM	
b. INTENDED USE(S) (Check Applicable Box(es))		
<input type="checkbox"/> GENERAL PURPOSE RADIOGRAPHY	<input type="checkbox"/> UROLOGY	<input type="checkbox"/> CT WHOLE BODY SCANNER
<input type="checkbox"/> GENERAL PURPOSE FLUOROSCOPY	<input type="checkbox"/> MAMMOGRAPHY	<input type="checkbox"/> HEAD - NECK (MEDICAL)
<input type="checkbox"/> TOMOGRAPHY (OTHER THAN CT)	<input type="checkbox"/> CHEST	<input type="checkbox"/> DENTAL - INTRAOURAL
<input type="checkbox"/> ANGIOGRAPHY	<input type="checkbox"/> CHIROPRACTIC	<input type="checkbox"/> DENTAL - CEPHALOMETRIC
<input type="checkbox"/> PODIATRY	<input type="checkbox"/> CT HEAD SCANNER	<input type="checkbox"/> DENTAL PANORAMIC
<input type="checkbox"/> VETERINARIAN	<input type="checkbox"/> RADIATION THERAPY SIMULATOR	
c. THE X-RAY SYSTEM IS (Check one)		e. DATE OF ASSEMBLY
<input type="checkbox"/> STATIONARY	<input type="checkbox"/> PORTABLE (HANDHELD)	____/____/____ (MM) (DD) (YYYY)
<input type="checkbox"/> MOBILE		d. STATE OF COLO. BLUE LABEL NUMBER (IF ON MACHINE)

4. COMPONENT INFORMATION

IF THIS IS THE REPLACEMENT OF A COMPONENT, INDICATE OLD MODEL & SERIAL NUMBER AND BLUE CDPHE CERTIFICATION LABEL NUMBER IN COMMENTS BELOW

a. THE MASTER CONTROL IS <input type="checkbox"/> A NEW INSTALLATION <input type="checkbox"/> EXISTING (Certified) <input type="checkbox"/> EXISTING (Non-certified)	b. CONTROL MANUFACTURER	d. CONTROL SERIAL NUMBER	e. DATE MANUFACTURED
c. CONTROL MODEL NUMBER		f. SYSTEM MODEL NAME (CT Systems Only)	

Complete the following information for the certified components listed below which you installed. For beam limiting devices, tables and CT gantries enter the manufacturer and Model number in the indicated spaces. For other certified components, enter in the appropriate blocks how many of each you installed in this system.

g. SELECTED COMPONENTS				h. OTHER CERTIFIED COMPONENTS (Enter number of each installed in appropriate blocks)									
BEAM LIMITING DEVICES	MANUFACTURER	MODEL NUMBER	DATE MANUFACTURED	<input type="checkbox"/> X-RAY CONTROL	<input type="checkbox"/> CRADLE								
	MANUFACTURER	MODEL NUMBER	DATE MANUFACTURED			<input type="checkbox"/> HIGH VOLTAGE GENERATOR	<input type="checkbox"/> FILM CHANGER						
TABLES	MANUFACTURER	MODEL NUMBER	DATE MANUFACTURED					<input type="checkbox"/> VERTICAL CASSETE HOLDER	<input type="checkbox"/> IMAGE INTENSIFIER				
	MANUFACTURER	MODEL NUMBER	DATE MANUFACTURED							<input type="checkbox"/> TUBE HOUSING ASSEMBLY	<input type="checkbox"/> SPOT FILM DEVICE		
CT GANTRY	MANUFACTURER	MODEL NUMBER	DATE MANUFACTURED									<input type="checkbox"/> DENTAL TUBE HEAD	<input type="checkbox"/> OTHER (Specify)
	MANUFACTURER	MODEL NUMBER	DATE MANUFACTURED										

5. ASSEMBLER CERTIFICATION

I affirm that all certified components assembled or installed by me, for which this report is being made, were adjusted and tested by me according to the instructions provided by the manufacture(s), were of the type required by the manufacturer(s), were of the type require by the diagnostic x-ray performance standard (21 CFR Part 1020), were not modified to adversely affect performance, and were installed in accordance with provisions of 21 CFR Part 1020. I also affirm that all instruction manuals and other information required by 21 CFR Part 1020 for this assembly and a copy of this form have been furnished to the purchaser and within 15 days from the date of assembly, a copy of this report will be distributed to the Colorado Department of Public Health and Environment, X-Ray Certification Unit.

a. PRINTED NAME	b. SIGNATURE	c. DATE
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6. COMMENTS (NOTE: Colorado Regulations require that the Assembler verifies the facility has a Shielding Design completed (when required) and the component(s) were installed according to the area diagram approved in the Shielding Design) SHIELDING DESIGN VERIFIED: YES NO N/A

7. DOES THE WORK YOU PERFORMED TODAY REQUIRE AN EVALUATION BY A CERTIFIED INSPECTOR? YES NO

PLEASE LEAVE COPY OF FORM WITH FACILITY

FORM CAN BE SUBMITTED VIA HTTP://CDPHE- CDPHE_XRAY_QISC@STATE.CO.US

FORM CDPHE 2579 (05/2018) PREVIOUS EDITION IS OBSOLETE