

FOR OFFICE USE ONLY (DATE RECEIVED)	COLORADO DEPARTMENT OF PUBLIC HEALTH & ENVIRONMENT HMWMD - X-Ray Certification Program 4300 Cherry Creek Drive South, B2 DENVER, COLORADO 80246-1530 (303) 692-3448 or 3443 FAX (303) 759-5355 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"/> REASSEMBLY - MIXED SYSTEM (Both certified and uncertified components)		<input type="checkbox"/> REPLACEMENT COMPONENTS IN AN EXISTING SYSTEM		<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 - INTRAORAL	
<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"/> RADIATION THERAPY SIMULATOR		<input type="checkbox"/> C-ARM FLUOROSCOPIC		<input type="checkbox"/> DIGITAL	
<input type="checkbox"/> BONE MINERAL ANALYSIS		<input type="checkbox"/> OTHER (Specify in Comments)			
c. THE X-RAY SYSTEM IS (Check one)			d. THE MASTER CONTROL IS IN ROOM		e. DATE OF ASSEMBLY
<input type="checkbox"/> STATIONARY					____/____/____
<input type="checkbox"/> MOBILE					(MM) (DD) (YYYY)

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

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 <input type="checkbox"/> HIGH VOLTAGE GENERATOR <input type="checkbox"/> FILM CHANGER <input type="checkbox"/> VERTICAL CASSETTE HOLDER <input type="checkbox"/> IMAGE INTENSIFIER <input type="checkbox"/> TUBE HOUSING ASSEMBLY <input type="checkbox"/> SPOT FILM DEVICE <input type="checkbox"/> DENTAL TUBE HEAD <input type="checkbox"/> OTHER (Specify)
	MANUFACTURER		MODEL NUMBER		DATE MANUFACTURED		
TABLES	MANUFACTURER		MODEL NUMBER		DATE MANUFACTURED		
	MANUFACTURER		MODEL NUMBER		DATE MANUFACTURED		
CT GANTRY	MANUFACTURER		MODEL NUMBER		DATE MANUFACTURED		
	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
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		
BLUE CDPHE CERTIFICATION LABEL NO.:		