

	DEPARTMENT OF HEALTH AND HUMAN SERVICES FOOD AND DRUG ADMINISTRATION REPORT OF ASSEMBLY OF A DIAGNOSTIC X-RAY SYSTEM	Form Approved: OMB No. 0910-0025 Expiration Date: February 28, 2026 See Reverse for PRA statement Assembler/Purchaser Control Number
Distribution List: Purchaser Assembler State Radiation Health Office		

1. EQUIPMENT LOCATION	2. ASSEMBLER INFORMATION
a. NAME OF HOSPITAL, DOCTOR OR OFFICE WHERE INSTALLED	a. COMPANY NAME
b. STREET ADDRESS	b. STREET ADDRESS
c. CITY	c. CITY
d. STATE	d. STATE
e. ZIP CODE	e. ZIP CODE
f. TELEPHONE NUMBER	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"/> REASSEMBLY - FULLY CERTIFIED SYSTEM	<input type="checkbox"/> REASSEMBLY - MIXED SYSTEM (Both certified and non-certified components)	<input type="checkbox"/> REPLACEMENT COMPONENTS IN AN EXISTING SYSTEM	
		<input type="checkbox"/> AN ADDITION TO AN EXISTING SYSTEM		
b. INTENDED USE(S) (Check appropriate box(es))				
<input type="checkbox"/> GENERAL PURPOSE RADIOGRAPHY	<input type="checkbox"/> UROLOGY	<input type="checkbox"/> CT WHOLE BODY SCANNER	<input type="checkbox"/> RADIATION THERAPY SIMULATOR	<input type="checkbox"/> OTHER (Specify in comments)
<input type="checkbox"/> GENERAL PURPOSE FLUOROSCOPY	<input type="checkbox"/> MAMMOGRAPHY	<input type="checkbox"/> HEAD-NECK (Medical)	<input type="checkbox"/> C-ARM FLUOROSCOPIC	
<input type="checkbox"/> TOMOGRAPHY (Other than CT)	<input type="checkbox"/> CHEST	<input type="checkbox"/> DENTAL-INTRAORAL	<input type="checkbox"/> DIGITAL	
<input type="checkbox"/> ANGIOGRAPHY	<input type="checkbox"/> CHIROPRACTIC	<input type="checkbox"/> DENTAL-CEPHALOMETRIC	<input type="checkbox"/> BONE MINERAL ANALYSIS	
<input type="checkbox"/> PODIATRY	<input type="checkbox"/> CT HEADSCANNER	<input type="checkbox"/> DENTAL PANORAMIC	<input type="checkbox"/> DENTAL-CT	
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			<div><div></div><div></div><div></div></div> <div>(mm) (dd) (yyyy)</div>	

4. COMPONENT INFORMATION (If additional space is needed for this section use another form, replacing the preprinted number with this Form Number, and complete Items 1, 4, and 5 only)

a. THE MASTER CONTROL IS	b. CONTROL MANUFACTURER	d. CONTROL SERIAL NUMBER	e. DATE MANUFACTURED
<input type="checkbox"/> A NEW INSTALLATION			
<input type="checkbox"/> EXISTING (Certified)	c. CONTROL MODEL NUMBER	f. SYSTEM MODEL NAME (CT Systems Only)	
<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			
	MANUFACTURER	MODEL NUMBER	DATE MANUFACTURED
	MANUFACTURER	MODEL NUMBER	DATE MANUFACTURED
	MANUFACTURER	MODEL NUMBER	DATE MANUFACTURED
	MANUFACTURER	MODEL NUMBER	DATE MANUFACTURED
	MANUFACTURER	MODEL NUMBER	DATE MANUFACTURED
h. OTHER CERTIFIED COMPONENTS (Enter number of each installed in appropriate blocks.)			
X-RAY CONTROL		CRADLE	
HIGH VOLTAGE GENERATOR		FILM CHANGER	
VERTICAL CASSETTE HOLDER		IMAGE INTENSIFIER	
TUBE HOUSING ASSEMBLY		SPOT FILM DEVICE	
DENTAL TUBE HEAD		FLUOROSCOPIC IMAGING ASSEMBLY	
CEPHALOMETRIC DEVICE		IMAGE RECEPTOR	
IMAGE RECEPTOR SUPPORT DEVICE		FLUOROSCOPIC AIR KERMA DISPLAY DEVICE	
OTHER			

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 required 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 have been furnished to the purchaser and, within 15 days following completion of the assembly, a copy of this form will be submitted to the purchaser and, where applicable, to the State agency responsible for radiation protection.	
a. PRINTED NAME	b. SIGNATURE

6. COMMENTS

Contact Information for State Radiation Health Offices is available on the website of the Conference of Radiation Control Program Directors (CRCPD),

<https://www.crcpd.org/mpage/Map>

Form may be downloaded at: <https://www.fda.gov/media/144454/download>

This section applies only to requirements of the Paperwork Reduction Act of 1995.

DO NOT SEND YOUR COMPLETED FORM TO THE PRA STAFF EMAIL ADDRESS BELOW.

The burden time for this collection of information is estimated to average 18 minutes per response, including the time to review instructions, search existing data sources, gather and maintain the data needed and complete and review the collection of information. Send comments regarding this burden estimate or any other aspect of this information collection, including suggestions for reducing this burden, to:

Department of Health and Human Services
Food and Drug Administration
Office of Chief Information Officer
Paperwork Reduction Act (PRA) Staff
PRASStaff@fda.hhs.gov

"An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB number."