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The document discusses FDA form, which reports the assembly of diagnostic Fillable FormReport of Assembly of a Diagnostic X-Ray System; Searchable FDA Forms Listing Federal regulations (SectionCFR (d)) require that an assembler who installs one or more certified components of a diagnostic x-ray system submit a report of assembly (Form FDA Complete the following information for the certified components listed below which you installed. REPORT Complete the following information for the certified components listed below which you installed. DEPARTMENT OF HEALTH AND HUMAN SERVICES. FDA Form Author: FDA Subject: Report of Assembly of a Diagnostic X-Ray System Created Date/17/ AM White OriginalFDA, Document Mail Center – WO, New Hampshire Avenue, Silver Spring, MD FORM FDA (09/20) PREVIOUS Form FDA FOR FDA USE ONLY. 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 3 Report of Assembly (Form FDA) is required for diagnostic x-ray components; see § (d)(1)-(3) of this chapterSystems records and reports are required if a manufacturer exercises the option and certifies the system as permitted in § (c) of this chapter This document supersedes Assembler's guide to diagnostic x-ray equipment: responsibilities of assemblers, distributors, and dealers of diagnostic x-ray equipment under the federal performance Public Health Service. For beam limiting devices, tables and CT gantries enter the manufacturer and FDA Form PDFFree download as PDF File.pdf), Text File.txt) or read online for free. FOOD AND DRUG ADMINISTRATION.

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