

**K970293 FUJI COMPUTED RADIOGRAPHY FCR/DICOM  
GATEWAY UNIT CR-DM666**

Mar 21, 1997  
56 days to decision

K970293 · Product code: **LMD** · Radiology  
Source: <https://www.510kdatabase.net/k970293/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Digital Image Communications, Radiological (LMD)
Date received	Jan 24, 1997
Decision date	Mar 21, 1997
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Fujifilm Medical System U.S.A., Inc.</b>
Location	Stamford, CT, US
Contact	ROBERT A UZENOFF
510(k) history	71 submissions · 71 cleared · 1988-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k970293/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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