

**K950610 FIKO FO**Apr 25, 1995  
74 days to decisionK950610 · Product code: **LMC** · Radiology  
Source: <https://www.510kdatabase.net/k950610/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Multi Format, Radiological (LMC)
Date received	Feb 10, 1995
Decision date	Apr 25, 1995
Days to decision	74 days
Third-party review	No
Summary / Statement	Summary
Other names	, TJER,A; O,AGER FTO-1000

**APPLICANT**

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Company	<b>Fujifilm Medical System U.S.A., Inc.</b>
Location	Stamford, CT, US
Contact	OBERT 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/k950610/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 16, 2026