

K840134 CAMTEX 1000Mar 16, 1984
72 days to decisionK840134 · Product code: **JAA** · Radiology
Source: <https://www.510kdatabase.net/k840134/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Fluoroscopic, Image-intensified (JAA)
Date received	Jan 4, 1984
Decision date	Mar 16, 1984
Days to decision	72 days
Third-party review	No

APPLICANT

Company	Digital Imaging Co. of America
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1984-1984

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Device record: <https://www.510kdatabase.net/k840134/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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