

**K811676 MAXI CAMERA/D**Jul 2, 1981  
17 days to decisionK811676 · Product code: **IYX** · Radiology  
Source: <https://www.510kdatabase.net/k811676/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Scintillation (gamma) (IYX)
Date received	Jun 15, 1981
Decision date	Jul 2, 1981
Days to decision	17 days
Third-party review	No

**APPLICANT**

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Company	<b>General Electric Co.</b>
Location	Mchenry, IL, US
510(k) history	254 submissions · 254 cleared · 1976-2011

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

Device record: <https://www.510kdatabase.net/k811676/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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