

**K812915 MX 125**Nov 24, 1981  
36 days to decisionK812915 · Product code: **ITY** · Radiology  
Source: <https://www.510kdatabase.net/k812915/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Assembly, Tube Housing, X-ray, Diagnostic (ITY)
Date received	Oct 19, 1981
Decision date	Nov 24, 1981
Days to decision	36 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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Device record: <https://www.510kdatabase.net/k812915/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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