

**K791830 L/U ARM - NEURO - VASCULAR #B5080B**Oct 26, 1979  
39 days to decisionK791830 · Product code: **JAA** · Radiology  
Source: <https://www.510kdatabase.net/k791830/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Fluoroscopic, Image-intensified (JAA)
Date received	Sep 17, 1979
Decision date	Oct 26, 1979
Days to decision	39 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/k791830/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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