

**K883664 LUM VASCULAR PROCEDURE SYSTEM #B5200A**Nov 7, 1988  
73 days to decisionK883664 · Product code: **IYB** · Radiology  
Source: <https://www.510kdatabase.net/k883664/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube Mount, X-ray, Diagnostic (IYB)
Date received	Aug 26, 1988
Decision date	Nov 7, 1988
Days to decision	73 days
Third-party review	No

**APPLICANT**

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Company	<b>General Electric Co.</b>
Location	Mchenry, IL, US
Contact	KROGER, PHD
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/k883664/>; 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 17, 2026