FDA 510(k) Premarket Notification

K883664 LUM VASCULAR PROCEDURE SYSTEM #B5200A

CLEARED

Nov 7, 1988
73 days to decision

K883664 · Product code: **IYB** · Radiology
Source: https://www.510kdatabase.net/k883664/

SUBMISSION DETAILS

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

Company	**General Electric Co.**
Location	Mchenry, IL, US
Contact	KROGER, PHD
510(k) history	254 submissions · 254 cleared · 1976-2011