FDA 510(k) Premarket Notification

K791830 L/U ARM - NEURO - VASCULAR #B5080B

CLEARED

Oct 26, 1979
39 days to decision

K791830 · Product code: **JAA** · Radiology
Source: https://www.510kdatabase.net/k791830/

SUBMISSION DETAILS

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

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