FDA 510(k) Premarket Notification

K811676 MAXI CAMERA/D

CLEARED

Jul 2, 1981
17 days to decision

K811676 · Product code: **IYX** · Radiology
Source: https://www.510kdatabase.net/k811676/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Scintillation (gamma) (IYX)
Date received	Jun 15, 1981
Decision date	Jul 2, 1981
Days to decision	17 days
Third-party review	No

APPLICANT

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