FDA 510(k) Premarket Notification

K801653 MAXI CAMERA 37

CLEARED

Aug 20, 1980
33 days to decision

K801653 · Product code: **IYX** · Radiology
Source: https://www.510kdatabase.net/k801653/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Scintillation (gamma) (IYX)
Date received	Jul 18, 1980
Decision date	Aug 20, 1980
Days to decision	33 days
Third-party review	No

APPLICANT

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