FDA 510(k) Premarket Notification

K812915 MX 125

CLEARED

Nov 24, 1981
36 days to decision

K812915 · Product code: **ITY** · Radiology
Source: https://www.510kdatabase.net/k812915/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Assembly, Tube Housing, X-ray, Diagnostic (ITY)
Date received	Oct 19, 1981
Decision date	Nov 24, 1981
Days to decision	36 days
Third-party review	No

APPLICANT

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