FDA 510(k) Premarket Notification

K811082 BETA I & BETA II

CLEARED

May 21, 1981
30 days to decision

K811082 · Product code: **IZL** · Radiology
Source: https://www.510kdatabase.net/k811082/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Mobile (IZL)
Date received	Apr 21, 1981
Decision date	May 21, 1981
Days to decision	30 days
Third-party review	No

APPLICANT

Company	**Porta Ray, Inc.**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1980-1986