

K811082 BETA I & BETA IIMay 21, 1981
30 days to decisionK811082 · 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

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Device record: <https://www.510kdatabase.net/k811082/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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