

K973833 CONTROL, X-RAY DIAGNOSTIC, GENERATOR, HIGH VOLTAGE, ASSEMBLY, TUBE HOUSING DIAGNOSTIC, BEAM LIMITING DEVICE, MANUAL

Dec 30, 1997
84 days to decision

K973833 · Product code: IZL · Radiology
Source: <https://www.510kdatabase.net/k973833/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Mobile (IZL)
Date received	Oct 7, 1997
Decision date	Dec 30, 1997
Days to decision	84 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Dynarad Corp.
Location	Deer Park, NY, US
Contact	RAYMOND MANEZ
510(k) history	9 submissions · 9 cleared · 1993-2000

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Device record: <https://www.510kdatabase.net/k973833/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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