

K833821 INTERAD 520Jan 13, 1984
72 days to decisionK833821 · Product code: **IYX** · Radiology
Source: <https://www.510kdatabase.net/k833821/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Scintillation (gamma) (IYX)
Date received	Nov 2, 1983
Decision date	Jan 13, 1984
Days to decision	72 days
Third-party review	No

APPLICANT

Company	Interad Systems, Inc.
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1981-1984

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Device record: <https://www.510kdatabase.net/k833821/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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