FDA 510(k) Premarket Notification

# K833821 INTERAD 520

**CLEARED**

Jan 13, 1984
72 days to decision

K833821 · Product code: **IYX** · Radiology
Source: https://www.510kdatabase.net/k833821/

## SUBMISSION DETAILS

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|---|---|
| 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 |