

**K874934 UNIVERSAL EXAMINATION TABLE**Jan 19, 1988  
46 days to decisionK874934 · Product code: **KXJ** · Radiology  
Source: <https://www.510kdatabase.net/k874934/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Radiologic (KXJ)
Date received	Dec 4, 1987
Decision date	Jan 19, 1988
Days to decision	46 days
Third-party review	No

**APPLICANT**

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Company	<b>Imatron, Inc.</b>
Location	South San Fransisco, CA, US
Contact	J. A CODUTO
510(k) history	11 submissions · 11 cleared · 1983-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k874934/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 19, 2026