

K840135 CRISAMar 23, 1984
80 days to decisionK840135 · Product code: **ITY** · Radiology
Source: <https://www.510kdatabase.net/k840135/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Assembly, Tube Housing, X-ray, Diagnostic (ITY)
Date received	Jan 3, 1984
Decision date	Mar 23, 1984
Days to decision	80 days
Third-party review	No

APPLICANT

Company	X-Ray Tube Corp.
Location	Walker, MI, US
510(k) history	3 submissions · 3 cleared · 1983-1984

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Device record: <https://www.510kdatabase.net/k840135/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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