FDA 510(k) Premarket Notification

K840135 CRISA

CLEARED

Mar 23, 1984
80 days to decision

K840135 · 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