FDA 510(k) Premarket Notification

K960575 ADVANTX LCV+

CLEARED

May 7, 1996
85 days to decision

K960575 · Product code: **IZI** · Radiology
Source: https://www.510kdatabase.net/k960575/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Angiographic (IZI)
Date received	Feb 12, 1996
Decision date	May 7, 1996
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**GE Medical Systems**
Location	Milwaukee, WI, US
Contact	LARRY A KROGER, PH.D.
510(k) history	169 submissions · 166 cleared · 1989-2022