FDA 510(k) Premarket Notification

K992603 XRT 600 R/F, WRF 0600

CLEARED

Oct 25, 1999
83 days to decision

K992603 · Product code: **JAA** · Radiology
Source: https://www.510kdatabase.net/k992603/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Fluoroscopic, Image-intensified (JAA)
Date received	Aug 3, 1999
Decision date	Oct 25, 1999
Days to decision	83 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Wuestec Medical, Inc.**
Location	Mobile, AL, US
Contact	BRENDA S DAVIS
510(k) history	9 submissions · 9 cleared · 1993-1999