

**K992603 XRT 600 R/F, WRF 0600**Oct 25, 1999  
83 days to decisionK992603 · Product code: **JAA** · Radiology  
Source: <https://www.510kdatabase.net/k992603/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k992603/> 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