

**K071233 XPERT 40 SPECIMEN RADIOGRAPHY SYSTEM**Jun 20, 2007  
48 days to decisionK071233 · Product code: **MWP** · Radiology  
Source: <https://www.510kdatabase.net/k071233/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cabinet, X-ray System (MWP)
Date received	May 3, 2007
Decision date	Jun 20, 2007
Days to decision	48 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Kub Technologies, Inc.</b>
Location	Shelton, CT, US
Contact	JOSEPH M AZARY
510(k) history	10 submissions · 10 cleared · 2007-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k071233/> 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 26, 2026