

**K163289 syngo. CT Single Source Dual Energy (twin beam)**Feb 9, 2017  
79 days to decisionK163289 · Product code: **JAK** · Radiology  
Source: <https://www.510kdatabase.net/k163289/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Tomography, Computed (JAK)
Date received	Nov 22, 2016
Decision date	Feb 9, 2017
Days to decision	79 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Medical Solutions USA, Inc., Ultrasound DI</b>
Location	Mountain View, CA, US
Contact	Kimberly Mangum
510(k) history	11 submissions · 11 cleared · 2007-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k163289/> 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 15, 2026