

**K200990 VIDAvision**Aug 7, 2020  
114 days to decisionK200990 · Product code: **JAK** · Radiology  
Source: <https://www.510kdatabase.net/k200990/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Tomography, Computed (JAK)
Date received	Apr 15, 2020
Decision date	Aug 7, 2020
Days to decision	114 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vida Diagnostics, Inc.</b>
Location	Iowa City, IA, US
Contact	Sandra Stapleton
510(k) history	2 submissions · 2 cleared · 2008-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k200990/> 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 June 19, 2026