

**K081556 CADSTREAM, VERSION 5.0**Feb 2, 2009  
244 days to decisionK081556 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k081556/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jun 3, 2008
Decision date	Feb 2, 2009
Days to decision	244 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Confirma, Inc.</b>
Location	Kirkland, WA, US
Contact	BRENT LEWIS
510(k) history	9 submissions · 9 cleared · 2002-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k081556/> 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 17, 2026