FDA 510(k) Premarket Notification

K081556 CADSTREAM, VERSION 5.0

CLEARED

Feb 2, 2009
244 days to decision

K081556 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k081556/

SUBMISSION DETAILS

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

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