

K090223 ICAD BREAST INTERVENTIONAL PLANNING SOFTWARE, MODEL D70024

Mar 31, 2009
60 days to decision

K090223 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k090223/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jan 30, 2009
Decision date	Mar 31, 2009
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Icad, Inc.
Location	Beavercreek, OH, US
Contact	JOHN A DELUCIA
510(k) history	14 submissions · 14 cleared · 2004-2024

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Device record: <https://www.510kdatabase.net/k090223/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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