

**K062237 CADI, VERSION 4**Nov 20, 2006  
110 days to decisionK062237 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k062237/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Aug 2, 2006
Decision date	Nov 20, 2006
Days to decision	110 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Synca Marketing, Inc.</b>
Location	Le Gardeur, Quebec, CA
Contact	STEPHANE ARSENAULT
510(k) history	2 submissions · 2 cleared · 2006-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k062237/> 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 18, 2026