

**K071086 RESOLUTIONMD CARDIAC PRODUCT FAMILY,
MODEL: RELEASE 1.0**Jun 1, 2007
45 days to decisionK071086 · Product code: LLZ · Radiology
Source: <https://www.510kdatabase.net/k071086/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Apr 17, 2007
Decision date	Jun 1, 2007
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Calgary Scientific, Inc.
Location	Calgary, Alberta, CA
Contact	PIERRE LEMIRE
510(k) history	8 submissions · 8 cleared · 2006-2016

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k071086/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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