FDA 510(k) Premarket Notification

K050846 B-CAD SYSTEM, VERSION 1.0

CLEARED

May 26, 2005
52 days to decision

K050846 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k050846/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Apr 4, 2005
Decision date	May 26, 2005
Days to decision	52 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**The Medipattern Corporation**
Location	Toronto, On, CA
Contact	PATRICIA A MILBANK
510(k) history	2 submissions · 2 cleared · 2005-2011

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k050846/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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