

K063267 ISITE PACS, V 4. XNov 9, 2006
10 days to decisionK063267 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k063267/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Oct 30, 2006
Decision date	Nov 9, 2006
Days to decision	10 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Stentor Inc., A Philips Medical Systems Co.
Location	Brisbane, CA, US
Contact	DEANA WISEMAN
510(k) history	1 submissions · 1 cleared · 2006-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k063267/> Data sourced from FDA 510(k) public records (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. - Generated May 18, 2026