

K073176 KANTERON DICOM STATION (KDS), MODEL 3.0Mar 19, 2008
127 days to decisionK073176 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k073176/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Nov 13, 2007
Decision date	Mar 19, 2008
Days to decision	127 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Monica Veytia
Location	Miami, FL, US
Contact	MONICA VEYTIA
510(k) history	1 submissions · 1 cleared · 2008-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k073176/> 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