

K103720 SYNAPSE 3D LUNG AND ABDOMEN ANALYSISMar 16, 2011
85 days to decisionK103720 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k103720/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Dec 21, 2010
Decision date	Mar 16, 2011
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Fujifilm Medical Systems U.S.A, Inc.
Location	Stamford, CT, US
Contact	KIMERLY A SHARP
510(k) history	39 submissions · 39 cleared · 2005-2018

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k103720/> 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 16, 2026