

K212777 Collaboration LiveSep 24, 2021
23 days to decisionK212777 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k212777/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Sep 1, 2021
Decision date	Sep 24, 2021
Days to decision	23 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Philips Ultrasound, Inc.
Location	Santa Ana, CA, US
Contact	Courtney Nix
510(k) history	46 submissions · 46 cleared · 1985-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k212777/> 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 13, 2026