FDA 510(k) Premarket Notification

K212777 Collaboration Live

CLEARED

Sep 24, 2021
23 days to decision

K212777 · 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