FDA 510(k) Premarket Notification

K242329 CT Collaboration Live

CLEARED

Nov 18, 2024
104 days to decision

K242329 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k242329/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Aug 6, 2024
Decision date	Nov 18, 2024
Days to decision	104 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Philips Healthcare (Suzhou) Co., Ltd.**
Location	Suzhou Jiangsu, CN
Contact	Yael Curtz
510(k) history	17 submissions · 17 cleared · 2014-2026