

K242329 CT Collaboration LiveNov 18, 2024
104 days to decisionK242329 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k242329/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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