

K221501 syngo.via View&GOOct 14, 2022
144 days to decisionK221501 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k221501/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	May 23, 2022
Decision date	Oct 14, 2022
Days to decision	144 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Siemens Healthcare GmbH
Location	Erlangen, DE
Contact	Frederike Jakob
510(k) history	30 submissions · 30 cleared · 2016-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k221501/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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