FDA 510(k) Premarket Notification

# K221501 syngo.via View&GO

**CLEARED**

Oct 14, 2022
144 days to decision

K221501 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k221501/

## SUBMISSION DETAILS

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|---|---|
| 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

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