FDA 510(k) Premarket Notification

# K232136 syngo.CT Applications

**CLEARED**

Jan 4, 2024
170 days to decision

K232136 · Product code: **JAK** · Radiology
Source: https://www.510kdatabase.net/k232136/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | System, X-ray, Tomography, Computed (JAK) |
| Date received | Jul 18, 2023 |
| Decision date | Jan 4, 2024 |
| Days to decision | 170 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Siemens Medical Solutions USA, Inc.** |
| Location | Hoffman Estates, IL, US |
| Contact | Clayton Ginn |
| 510(k) history | 778 submissions · 778 cleared · 1980-2026 |