

**K191040 syngo.via (Version VB40A)**May 16, 2019  
27 days to decisionK191040 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k191040/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Apr 19, 2019
Decision date	May 16, 2019
Days to decision	27 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Healthcare GmbH</b>
Location	Erlangen, DE
Contact	Elango Alampalayam Rangappan
510(k) history	30 submissions · 30 cleared · 2016-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Tuv Sud America, Inc.</b>
Contact	Georg Bauer

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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