

**K192462 syngo.MR Applications**Jan 31, 2020  
144 days to decisionK192462 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k192462/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Sep 9, 2019
Decision date	Jan 31, 2020
Days to decision	144 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Medical Solutions USA, Inc.</b>
Location	Hoffman Estates, IL, US
Contact	Cordell L. Fields
510(k) history	779 submissions · 779 cleared · 1980-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k192462/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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