

**K182050 Linear Accelerator Control Console Software v13.1.010  
Update, syngo Radiation Therapy Therapist v4.3.1\_MR3 Patch,  
syngo Radiation Therapy Oncologist v4.3.1\_MR3 Patch**

Oct 3, 2018  
64 days to decision

K182050 · Product code: **IYE** · Radiology  
Source: <https://www.510kdatabase.net/k182050/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Jul 31, 2018
Decision date	Oct 3, 2018
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Medical Solutions USA, Inc.</b>
Location	Hoffman Estates, IL, US
Contact	Martin Rajchel
510(k) history	778 submissions · 778 cleared · 1980-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k182050/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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