

**K181613 MAGNETOM Avanto, MAGNETOM Verio with software syngo MR E11D**Nov 6, 2018  
140 days to decisionK181613 · Product code: LNH · Radiology  
Source: <https://www.510kdatabase.net/k181613/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jun 19, 2018
Decision date	Nov 6, 2018
Days to decision	140 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k181613/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](https://accessdata.fda.gov). - Generated May 15, 2026