

**K160523 BrightMatter Guide with BrightMatter Pointer**Jun 24, 2016  
120 days to decisionK160523 · Product code: **HAW** · Neurology  
Source: <https://www.510kdatabase.net/k160523/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Neurological Stereotaxic Instrument (HAW)
Date received	Feb 25, 2016
Decision date	Jun 24, 2016
Days to decision	120 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Synaptive Medical, Inc.</b>
Location	Toronto On, CA
Contact	Cameron Piron
510(k) history	10 submissions · 10 cleared · 2014-2024

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k160523/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 26, 2026