

**K964229 REGULUS NAVIGATOR**Aug 19, 1997  
300 days to decisionK964229 · Product code: **HAW** · Neurology  
Source: <https://www.510kdatabase.net/k964229/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Neurological Stereotaxic Instrument (HAW)
Date received	Oct 23, 1996
Decision date	Aug 19, 1997
Days to decision	300 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Compass Intl., Inc.</b>
Location	Rochester, MN, US
Contact	DEBRAH A FISHER
510(k) history	2 submissions · 2 cleared · 1995-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k964229/> 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 20, 2026