

**K062640 STRYKER NAVIGATION SYSTEM - CRANIAL MODULE,  
MODEL 6000-650-000**Dec 14, 2006  
99 days to decisionK062640 · Product code: **HAW** · Neurology  
Source: <https://www.510kdatabase.net/k062640/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Neurological Stereotaxic Instrument (HAW)
Date received	Sep 6, 2006
Decision date	Dec 14, 2006
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Instruments</b>
Location	Kalamazoo, MI, US
Contact	BECKEY DITTY
510(k) history	73 submissions · 73 cleared · 1994-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k062640/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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