

**K991256 VIEWPOINT ENT/ORTHOPEDIC OPTION**Oct 8, 1999  
178 days to decisionK991256 · Product code: **HAW** · Neurology  
Source: <https://www.510kdatabase.net/k991256/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Neurological Stereotaxic Instrument (HAW)
Date received	Apr 13, 1999
Decision date	Oct 8, 1999
Days to decision	178 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Philips Medical Systems (Cleveland), Inc.</b>
Location	Mchenry, IL, US
Contact	ELAINE K KEELER
510(k) history	190 submissions · 190 cleared · 1977-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k991256/> 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 16, 2026