

**K023623 NOVAVISION, MODEL 2.0**Apr 22, 2003  
175 days to decisionK023623 · Product code: **HPT** · Neurology  
Source: <https://www.510kdatabase.net/k023623/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Perimeter, Automatic, Ac-powered (HPT)
Date received	Oct 29, 2002
Decision date	Apr 22, 2003
Days to decision	175 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Novavision, Inc.</b>
Location	Washington, DC, US
Contact	SANDRA J.P. DENNIS
510(k) history	3 submissions · 3 cleared · 1996-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k023623/> 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 June 29, 2026