

**K020764 NAVIPRO**Jun 3, 2002  
88 days to decisionK020764 · Product code: **HAW** · Neurology  
Source: <https://www.510kdatabase.net/k020764/>**SUBMISSION DETAILS**

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|                       |   |
|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)        |
| Submission type       | Traditional                               |
| Device classification | Neurological Stereotaxic Instrument (HAW) |
| Date received         | Mar 7, 2002                               |
| Decision date         | Jun 3, 2002                               |
| Days to decision      | 88 days                                   |
| Third-party review    | No  |
| Summary / Statement   | Summary                                   |

**APPLICANT**

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|                |   |
|----------------|---|
| Company        | <b>Kinamed, Inc.</b>                    |
| Location       | Los Angeles, CA, US                     |
| Contact        | VINEET K SARIN                          |
| 510(k) history | 33 submissions · 29 cleared · 1988-2013 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020764/> 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 19, 2026