

**K953173 NONPOWERED NEUROSURGICAL INSTRUMENT**Oct 3, 1995  
88 days to decisionK953173 · Product code: **GWG** · Neurology  
Source: <https://www.510kdatabase.net/k953173/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Endoscope, Neurological (GWG)      |
| Date received         | Jul 7, 1995                        |
| Decision date         | Oct 3, 1995                        |
| Days to decision      | 88 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---|
| Company        | <b>United States Surgical, A Division of Tyco Healthc</b> |
| Location       | Mchenry, IL, US   |
| Contact        | JANET G JOHNSON   |
| 510(k) history | 218 submissions · 200 cleared · 1977-2007                 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k953173/> 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