

**K031787 SMITH & NEPHEW IUR MORCELLATION SYSTEM**Dec 2, 2003  
175 days to decisionK031787 · Product code: **HIH** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k031787/>**SUBMISSION DETAILS**

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|                       |                                      |
|-----------------------|--------------------------------------|
| Decision              | Substantially Equivalent (Cleared)   |
| Submission type       | Traditional                          |
| Device classification | Hysteroscope (and Accessories) (HIH) |
| Date received         | Jun 10, 2003                         |
| Decision date         | Dec 2, 2003                          |
| Days to decision      | 175 days                             |
| Third-party review    | No                                   |
| Summary / Statement   | Summary                              |

**APPLICANT**

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|----------------|---|
| Company        | <b>Smith &amp; Nephew, Inc.</b>   |
| Location       | Mchenry, IL, US   |
| Contact        | JANICE HASELTON   |
| Website        | <a href="http://www.smith-nephew.com/">http://www.smith-nephew.com/</a> |
| 510(k) history | 530 submissions · 517 cleared · 1980-2026                               |

Smith & Nephew, Inc. is a medical technology company focused on repair, regeneration, and replacement of soft and hard tissues. The company operates with a manufacturing facility in McHenry, US. Smith & Nephew has established a significant regulatory track record with the FDA. The company has received FDA 510(k) clearances from total submissions since 1980. Orthopedic devices represent the dominant category, accounting for 71% of submissions. The company remains active, with the latest clearance in 2025. Recent cleared devices reflect a strong focus on orthopedic surgical...

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