

**K041601 SURFIX KNEE OSTEOTOMY SYSTEM**Aug 27, 2004  
74 days to decisionK041601 · Product code: **HRS** · Orthopedic  
Source: <https://www.510kdatabase.net/k041601/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Plate, Fixation, Bone (HRS)        |
| Date received         | Jun 14, 2004                       |
| Decision date         | Aug 27, 2004                       |
| Days to decision      | 74 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Surfix Technologies</b>            |
| Location       | Potomac, MD, US                       |
| Contact        | NORMAN F ESTRIN                       |
| 510(k) history | 1 submissions · 1 cleared · 2004-2004 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k041601/> 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