

**K980637 IMPLEX CONTINUUM KNEE UNITY TIBIAL COMPONENT**Dec 16, 1998  
300 days to decisionK980637 · Product code: **JWH** · Orthopedic  
Source: <https://www.510kdatabase.net/k980637/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)   |
| Submission type       | Traditional  |
| Device classification | Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH) |
| Date received         | Feb 19, 1998   |
| Decision date         | Dec 16, 1998   |
| Days to decision      | 300 days   |
| Third-party review    | No   |
| Summary / Statement   | Summary  |

**APPLICANT**

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|----------------|---|
| Company        | <b>Implex Corp.</b>                     |
| Location       | Allendale, NJ, US                       |
| Contact        | ROBERT POGGIE, PH.D.                    |
| 510(k) history | 65 submissions · 61 cleared · 1993-2005 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k980637/>; Data sourced from FDA 510(k) public records (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 14, 2026