

**K971789 IMPLEX CONTINUUM POROUS PATELLAR**Jul 24, 1997  
71 days to decisionK971789 · Product code: **JWH** · Orthopedic  
Source: <https://www.510kdatabase.net/k971789/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	May 14, 1997
Decision date	Jul 24, 1997
Days to decision	71 days
Third-party review	No

**APPLICANT**

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Company	<b>Implex Corp.</b>
Location	Allendale, NJ, US
Contact	GLENN N BYRD
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/k971789/>; 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 14, 2026