

**K012095 THE LPS - CONTINUUM TM HYBRID TIBIA, MODELS
32-5886-2Y-XX (ZIMMER) AND 05-119-XXYY1-0 (IMPLEX)**Jul 26, 2001
21 days to decisionK012095 · Product code: **JWH** · Orthopedic
Source: <https://www.510kdatabase.net/k012095/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Jul 5, 2001
Decision date	Jul 26, 2001
Days to decision	21 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Implex Corp.
Location	Allendale, NJ, US
Contact	ROBERT A POGGIE
510(k) history	65 submissions · 61 cleared · 1993-2005

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k012095/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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