

**K192507 LOSPA II Knee System**Dec 11, 2019  
90 days to decisionK192507 · Product code: **JWH** · Orthopedic  
Source: <https://www.510kdatabase.net/k192507/>**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	Sep 12, 2019
Decision date	Dec 11, 2019
Days to decision	90 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Corentec Co., Ltd.</b>
Location	West Cadwell, NJ, US
Contact	J.S. Daniel
510(k) history	33 submissions · 33 cleared · 2010-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k192507/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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