

**K021908 PROVEN MODULAR POSTERIOR STABILIZED FEMORAL**

Jul 9, 2002  
29 days to decision

K021908 · Product code: **JWH** · Orthopedic  
Source: <https://www.510kdatabase.net/k021908/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Jun 10, 2002
Decision date	Jul 9, 2002
Days to decision	29 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Stelkast Company</b>
Location	Pittsburgh, PA, US
Contact	DONALD A STEVENS
510(k) history	40 submissions · 37 cleared · 1994-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k021908/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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