

**K982222 SYNTHES (USA) DISTAL FEMUR PLATE (DFP) SYSTEM**

Jul 29, 1998  
35 days to decision

K982222 · Product code: **HWC** · Orthopedic  
Source: <https://www.510kdatabase.net/k982222/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	Jun 24, 1998
Decision date	Jul 29, 1998
Days to decision	35 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Synthes (Usa)</b>
Location	Mchenry, IL, US
Contact	SHERI L MUSGNUNG
510(k) history	411 submissions · 394 cleared · 1977-2015

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

Device record: <https://www.510kdatabase.net/k982222/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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