

**K972393 INTER-OP HA POROUS ACETABULAR SYSTEM  
(HA/CSTI)**Sep 19, 1997  
85 days to decisionK972393 · Product code: **LPH** · Orthopedic  
Source: <https://www.510kdatabase.net/k972393/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Porous Uncemented (LPH)
Date received	Jun 26, 1997
Decision date	Sep 19, 1997
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sulzer Orthopedics, Inc.</b>
Location	Austin, TX, US
Contact	JACQUELYN HUGHES
510(k) history	45 submissions · 41 cleared · 1997-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k972393/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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