

**K935295 ALTA FEMORAL INTRAMEDULLARY RODS/CFX  
RODS MODIFICATION**Oct 11, 1994  
343 days to decisionK935295 · Product code: **HSB** · Orthopedic  
Source: <https://www.510kdatabase.net/k935295/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Rod, Fixation, Intramedullary And Accessories (HSB)
Date received	Nov 2, 1993
Decision date	Oct 11, 1994
Days to decision	343 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Howmedica Corp.</b>
Location	Mchenry, IL, US
Contact	ROBERT E SMITH
510(k) history	373 submissions · 325 cleared · 1976-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k935295/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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