

**K931678 ORTHOMET LHMC FEMORAL STEM**Oct 14, 1994  
556 days to decisionK931678 · Product code: **HMC** · Ophthalmic  
Source: <https://www.510kdatabase.net/k931678/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Eye Movement, Diagnostic (HMC)
Date received	Apr 6, 1993
Decision date	Oct 14, 1994
Days to decision	556 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Orthomet, Inc.</b>
Location	Plymouth, MN, US
Contact	DAVID A CANNISTRACI
510(k) history	60 submissions · 41 cleared · 1986-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k931678/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 21, 2026