

**K810540 KNEE PROSTHESIS**Apr 7, 1981  
39 days to decisionK810540 · Product code: **HRY** · Orthopedic  
Source: <https://www.510kdatabase.net/k810540/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Semi-constrained, Cemented, Metal/polymer (HRY)
Date received	Feb 27, 1981
Decision date	Apr 7, 1981
Days to decision	39 days
Third-party review	No

**APPLICANT**

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Company	<b>Cintor Orthopaedic Div.</b>
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1980-1981

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k810540/> 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 23, 2026