

**K020741 UNICONDYLAR KNEE**Jun 4, 2002  
90 days to decisionK020741 · Product code: **HRY** · Orthopedic  
Source: <https://www.510kdatabase.net/k020741/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Semi-constrained, Cemented, Metal/polymer (HRY)
Date received	Mar 6, 2002
Decision date	Jun 4, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Encore Orthopedics, Inc.</b>
Location	Red Rock, TX, US
Contact	JOANNA DROEGE
510(k) history	85 submissions · 74 cleared · 1993-2004

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020741/> 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 20, 2026