

**K051721 GLOBAL RESURFACING UNICOMPARTMENTAL
KNEE SYSTEM**Oct 26, 2005
121 days to decisionK051721 · Product code: **HRY** · Orthopedic
Source: <https://www.510kdatabase.net/k051721/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Semi-constrained, Cemented, Metal/polymer (HRY)
Date received	Jun 27, 2005
Decision date	Oct 26, 2005
Days to decision	121 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Global Orthopaedic Technology, USA, Inc.
Location	Oviedo, FL, US
Contact	CARL KNOBLOCH
510(k) history	3 submissions · 3 cleared · 2004-2005

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k051721/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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