FDA 510(k) Premarket Notification

K020741 UNICONDYLAR KNEE

CLEARED

Jun 4, 2002
90 days to decision

K020741 · 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	**Encore Orthopedics, Inc.**
Location	Red Rock, TX, US
Contact	JOANNA DROEGE
510(k) history	85 submissions · 74 cleared · 1993-2004