FDA 510(k) Premarket Notification

K022143 INTERPORE CROSS INTERNATIONAL ANTERIOR FIXATION DEVICE (AFD)

CLEARED

Jan 23, 2003
205 days to decision

K022143 · Product code: **MQP** · Orthopedic
Source: https://www.510kdatabase.net/k022143/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Jul 2, 2002
Decision date	Jan 23, 2003
Days to decision	205 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Interpore Cross Intl.**
Location	Irvine, CA, US
Contact	LYNN M RODARTI
510(k) history	39 submissions · 38 cleared · 1998-2005