

K043229 MODIFICATION TO ALTIUS OCT SYSTEMDec 21, 2004
29 days to decisionK043229 · Product code: **KWP** · Orthopedic
Source: <https://www.510kdatabase.net/k043229/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Nov 22, 2004
Decision date	Dec 21, 2004
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k043229/> Data sourced from FDA 510(k) public records (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. - Generated May 20, 2026