

K002592 INTERPORE CROSS ANTERIOR CERVICAL PLATE SYSTEM

Oct 6, 2000
46 days to decision

K002592 · Product code: **KWQ** · Orthopedic
Source: <https://www.510kdatabase.net/k002592/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Aug 21, 2000
Decision date	Oct 6, 2000
Days to decision	46 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k002592/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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