FDA 510(k) Premarket Notification

# K002592 INTERPORE CROSS ANTERIOR CERVICAL PLATE SYSTEM

**CLEARED**

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 |