FDA 510(k) Premarket Notification

# K974757 KANEDA ANTERIOR SCOLIOSIS SYSTEM KASS

**CLEARED**

Mar 5, 1998
76 days to decision

K974757 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k974757/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent - SN |
| Submission type | Traditional |
| Device classification | Appliance, Fixation, Spinal Intervertebral Body (KWQ) |
| Date received | Dec 19, 1997 |
| Decision date | Mar 5, 1998 |
| Days to decision | 76 days |
| Third-party review | No |

## APPLICANT

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|---|---|
| Company | **Acromed Corp.** |
| Location | Cleveland, OH, US |
| Contact | GREG CANNEDY |
| 510(k) history | 41 submissions · 22 cleared · 1984-1998 |