

**K974757 KANEDA ANTERIOR SCOLIOSIS SYSTEM KASS**Mar 5, 1998  
76 days to decisionK974757 · 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	<b>Acromed Corp.</b>
Location	Cleveland, OH, US
Contact	GREG CANNEDY
510(k) history	41 submissions · 22 cleared · 1984-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k974757/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 21, 2026