

**K970462 ACROMED ANTERIOR CERVICAL STABILIZATION SYSTEM**Apr 3, 1997  
56 days to decisionK970462 · Product code: **KWQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k970462/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Feb 6, 1997
Decision date	Apr 3, 1997
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Acromed Corp.</b>
Location	Cleveland, OH, US
Contact	SLLY LEVINE
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/k970462/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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