

K895439 ACROMED BEADED SUBLAMINAR WIRENov 22, 1989
78 days to decisionK895439 · Product code: **KWP** · Orthopedic
Source: <https://www.510kdatabase.net/k895439/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Sep 5, 1989
Decision date	Nov 22, 1989
Days to decision	78 days
Third-party review	No

APPLICANT

Company	Acromed Corp.
Location	Cleveland, OH, US
Contact	ZANG, RN
510(k) history	41 submissions · 22 cleared · 1984-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k895439/>; Data sourced from FDA 510(k) public records (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. - Generated May 21, 2026