

K092124 AXIALIF 2L, AXIALIF 2-LEVEL SYSTEM, AXIALIF APOLLO, AXIALIF IIJan 21, 2010
190 days to decisionK092124 · Product code: **KWQ** · Orthopedic
Source: <https://www.510kdatabase.net/k092124/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Jul 15, 2009
Decision date	Jan 21, 2010
Days to decision	190 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Trans1 Incorporated
Location	Wilmington, NC, US
Contact	CHERYL WAGONER
510(k) history	9 submissions · 9 cleared · 2005-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k092124/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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