

K951268 SPINE-TECH MODIFIED LAPAROSCOPIC SPINAL FUSION INSTRUMENT SYSTEMJul 31, 1995
132 days to decisionK951268 · Product code: **HRX** · Orthopedic
Source: <https://www.510kdatabase.net/k951268/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Mar 21, 1995
Decision date	Jul 31, 1995
Days to decision	132 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Spine-Tech, Inc.
Location	Minneapolis, MN, US
Contact	DAVID A CANNISTRACI
510(k) history	4 submissions · 3 cleared · 1994-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k951268/> 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