

K946258 MODEL 8000 ATLAS C-1 ORTHOGONAL ADJUSTING INSTRUMENTNov 20, 1995
349 days to decisionK946258 · Product code: **LXM** · Physical Medicine
Source: <https://www.510kdatabase.net/k946258/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Manipulator, Plunger-like Joint (LXM)
Date received	Dec 6, 1994
Decision date	Nov 20, 1995
Days to decision	349 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Spinalight, Inc.
Location	Woodstock, GA, US
Contact	T A COX
510(k) history	7 submissions · 7 cleared · 1994-2018

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k946258/> 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 30, 2026