

**K062670 VERTIFLEX SPINAL SCREW SYSTEM**Jan 12, 2007  
127 days to decisionK062670 · Product code: **MNH** · Orthopedic  
Source: <https://www.510kdatabase.net/k062670/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spondylolisthesis Spinal Fixation (MNH)
Date received	Sep 7, 2006
Decision date	Jan 12, 2007
Days to decision	127 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Vertiflex (Tm), Incorporated</b>
Location	Carlsbad, CA, US
Contact	STEVE REITZLER
510(k) history	2 submissions · 2 cleared · 2007-2007

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k062670/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 19, 2026