

K092076 AEGIS & AEGIS II SPINAL SYSTEMApr 2, 2010
267 days to decisionK092076 · Product code: **MNI** · Orthopedic
Source: <https://www.510kdatabase.net/k092076/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spinal Pedicle Fixation (MNI)
Date received	Jul 9, 2009
Decision date	Apr 2, 2010
Days to decision	267 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Corentec Co., Ltd.
Location	West Cadwell, NJ, US
Contact	ROBERT SCHIFF
510(k) history	33 submissions · 33 cleared · 2010-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k092076/> 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 18, 2026