

**K003928 TOWNLEY TRANSFACETPEDICULAR SCREW
FIXATION SYSTEM**Jan 17, 2001
28 days to decisionK003928 · Product code: **MRW** · Orthopedic
Source: <https://www.510kdatabase.net/k003928/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Facet Screw Spinal Device (MRW)
Date received	Dec 20, 2000
Decision date	Jan 17, 2001
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Sofamor Danek, Inc.
Location	Memphis, TN, US
Contact	RICHARD W TREHARNE
510(k) history	99 submissions · 89 cleared · 2000-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k003928/> 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 17, 2026