

**K013829 MODIFICATION TO TOWNLEY
TRANSFACETPEDICULAR SCREW FIXATION SYSTEM**Dec 19, 2001
30 days to decisionK013829 · Product code: **MRW** · Orthopedic
Source: <https://www.510kdatabase.net/k013829/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Facet Screw Spinal Device (MRW)
Date received	Nov 19, 2001
Decision date	Dec 19, 2001
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k013829/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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