FDA 510(k) Premarket Notification

K001153 MODIFICATION TO STEATHSTATION TREATMENT GUIDANCE PLATFORM

CLEARED

May 3, 2000
23 days to decision

K001153 · Product code: **HAW** · Neurology
Source: https://www.510kdatabase.net/k001153/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Neurological Stereotaxic Instrument (HAW)
Date received	Apr 10, 2000
Decision date	May 3, 2000
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medtronic Surgical Navigation**
Location	Broomfield, CO, US
Contact	VICTORIA RENDON
510(k) history	3 submissions · 3 cleared · 2000-2005

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k001153/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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