

**K083371 NAVIGATOR PRO WITH AEP SOFTWARE (BIOMARK,
ENOG, CHAMP)**Dec 8, 2009
389 days to decisionK083371 · Product code: **GWF** · Neurology
Source: <https://www.510kdatabase.net/k083371/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Nov 14, 2008
Decision date	Dec 8, 2009
Days to decision	389 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Natus Medical, Inc.
Location	San Carlos, CA, US
Contact	NICOHL R WILDING
510(k) history	12 submissions · 12 cleared · 1995-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k083371/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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