FDA 510(k) Premarket Notification

K871338 NOVAMETRIX MODEL 200 MAGSTIM

CLEARED

Aug 4, 1987
123 days to decision

K871338 · Product code: **GWF** · Neurology
Source: https://www.510kdatabase.net/k871338/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Apr 3, 1987
Decision date	Aug 4, 1987
Days to decision	123 days
Third-party review	No

APPLICANT

Company	**Novametrix Medical Systems, Inc.**
Location	Mchenry, IL, US
Contact	JAMES R VEALE
510(k) history	45 submissions · 45 cleared · 1978-2001