FDA 510(k) Premarket Notification

K931923 DANTEC MAGLITE MAGNETIC STIMULATOR (MAGLITE)

CLEARED

Nov 9, 1993
207 days to decision

K931923 · Product code: **GWF** · Neurology
Source: https://www.510kdatabase.net/k931923/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Apr 16, 1993
Decision date	Nov 9, 1993
Days to decision	207 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Dantec Medical, Inc.**
Location	Mahwah, NJ, US
Contact	RICHARD D MANTHEI
510(k) history	25 submissions · 25 cleared · 1990-1996