FDA 510(k) Premarket Notification

K870567 EVOPORT 300

CLEARED

Oct 6, 1987
238 days to decision

K870567 · Product code: **GWF** · Neurology
Source: https://www.510kdatabase.net/k870567/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Feb 10, 1987
Decision date	Oct 6, 1987
Days to decision	238 days
Third-party review	No

APPLICANT

Company	**Dantec Electronics, Inc.**
Location	Walker, MI, US
Contact	HENRIK HENRIKSEN
510(k) history	12 submissions · 12 cleared · 1985-1990