FDA 510(k) Premarket Notification

K833037 POINTER F-3

CLEARED

Nov 29, 1983
83 days to decision

K833037 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k833037/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Sep 7, 1983
Decision date	Nov 29, 1983
Days to decision	83 days
Third-party review	No

APPLICANT

Company	**Petra Intl. Co.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983