FDA 510(k) Premarket Notification

K833354 AUTO-TENS MODEL VI DUAL CHANNEL BATT

CLEARED

Nov 28, 1983
61 days to decision

K833354 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k833354/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Sep 28, 1983
Decision date	Nov 28, 1983
Days to decision	61 days
Third-party review	No

APPLICANT

Company	**Physio Tronics Corp.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983