FDA 510(k) Premarket Notification

K880042 ACUTRON

CLEARED

Aug 1, 1988
209 days to decision

K880042 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k880042/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jan 5, 1988
Decision date	Aug 1, 1988
Days to decision	209 days
Third-party review	No

APPLICANT

Company	**Davis Assoc.**
Location	Bridgeport, CT, US
Contact	PAUL DAVIS
510(k) history	1 submissions · 1 cleared · 1988-1988