FDA 510(k) Premarket Notification

K811939 VITATRON 88

CLEARED

Jul 23, 1981
17 days to decision

K811939 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k811939/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jul 6, 1981
Decision date	Jul 23, 1981
Days to decision	17 days
Third-party review	No

APPLICANT

Company	**Vitatronics**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981