FDA 510(k) Premarket Notification

K832296 S2 POSTERIOR-TIB

CLEARED

Aug 31, 1983
49 days to decision

K832296 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k832296/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Jul 13, 1983
Decision date	Aug 31, 1983
Days to decision	49 days
Third-party review	No

APPLICANT

Company	**Anstek Corp.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1983-1983