FDA 510(k) Premarket Notification

K820082 POST-OP ELECTRODE (TENS ELECTRODE)

CLEARED

Jan 28, 1982
16 days to decision

K820082 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k820082/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Jan 12, 1982
Decision date	Jan 28, 1982
Days to decision	16 days
Third-party review	No

APPLICANT

Company	**Consolidated Medical Equipment, Inc.**
Location	Mchenry, IL, US
510(k) history	15 submissions · 15 cleared · 1980-1986