FDA 510(k) Premarket Notification

K960885 MODEL 1240, SMALL E NET

CLEARED

Apr 26, 1996
53 days to decision

K960885 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k960885/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Mar 4, 1996
Decision date	Apr 26, 1996
Days to decision	53 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Physiometrix, Inc.**
Location	Sunnyvale, CA, US
Contact	DAWN E FRAZER
510(k) history	22 submissions · 22 cleared · 1992-2005