FDA 510(k) Premarket Notification

K933807 GROUND PLATE ELECTRODE

CLEARED

Jul 26, 1994
356 days to decision

K933807 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k933807/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Aug 4, 1993
Decision date	Jul 26, 1994
Days to decision	356 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cadwell Laboratories, Inc.**
Location	Walker, MI, US
Contact	CARLTON M CADWELL
510(k) history	46 submissions · 46 cleared · 1979-2007