FDA 510(k) Premarket Notification

K963307 GDC PATIENT RETURN ELECTTRODE MODEL 45021

CLEARED

Dec 17, 1996
117 days to decision

K963307 · Product code: **HCG** · Neurology
Source: https://www.510kdatabase.net/k963307/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Neurovascular Embolization (HCG)
Date received	Aug 22, 1996
Decision date	Dec 17, 1996
Days to decision	117 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Target Therapeutics**
Location	Los Angeles, CA, US
Contact	LARAINE PANGELINA
510(k) history	70 submissions · 70 cleared · 1985-1998