FDA 510(k) Premarket Notification

K960762 SYNERGY PLUS

CLEARED

Jun 13, 1996
108 days to decision

K960762 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k960762/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Feb 26, 1996
Decision date	Jun 13, 1996
Days to decision	108 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**The Prometheus Group**
Location	Portsmouth, NH, US
Contact	RICHARD M HORTON
510(k) history	11 submissions · 11 cleared · 1991-2016