

K960762 SYNERGY PLUSJun 13, 1996
108 days to decisionK960762 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k960762/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 16, 2026