FDA 510(k) Premarket Notification

K842804 DIADYN 4

CLEARED

Aug 21, 1984
35 days to decision

K842804 · Product code: **IPF** · Physical Medicine
Source: https://www.510kdatabase.net/k842804/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Jul 17, 1984
Decision date	Aug 21, 1984
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**Robert Bosch Corp.**
Location	Mchenry, IL, US
510(k) history	21 submissions · 18 cleared · 1979-1986