

K842804 DIADYN 4Aug 21, 1984
35 days to decisionK842804 · Product code: **IPF** · Physical MedicineSource: <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

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Device record: <https://www.510kdatabase.net/k842804/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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