FDA 510(k) Premarket Notification

K913898 MYOTRAC+/S, MYODAC+/S

CLEARED

Mar 26, 1992
209 days to decision

K913898 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k913898/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Aug 30, 1991
Decision date	Mar 26, 1992
Days to decision	209 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Thought Technology , Ltd.**
Location	Mchenry, IL, US
Contact	HAL K MYERS
510(k) history	22 submissions · 22 cleared · 1981-2022