

**K913898 MYOTRAC+ /S, MYODAC+ /S**Mar 26, 1992  
209 days to decisionK913898 · 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	<b>Thought Technology , Ltd.</b>
Location	Mchenry, IL, US
Contact	HAL K MYERS
510(k) history	22 submissions · 22 cleared · 1981-2022

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k913898/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 14, 2026