

K761350 MYOTRONMar 15, 1977
77 days to decisionK761350 · Product code: **HCC** · Neurology
Source: <https://www.510kdatabase.net/k761350/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Dec 28, 1976
Decision date	Mar 15, 1977
Days to decision	77 days
Third-party review	No

APPLICANT

Company	Somatronics, Inc.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1977-1979

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Device record: <https://www.510kdatabase.net/k761350/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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