

**K970051 MUSCLESense II CLINICAL ELECTROMYOGRAPHY
DEVICE**Jul 10, 1997
184 days to decisionK970051 · Product code: **HCC** · Neurology
Source: <https://www.510kdatabase.net/k970051/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Jan 7, 1997
Decision date	Jul 10, 1997
Days to decision	184 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Smith and Nephew Donjoy, Inc.
Location	Carsbad, CA, US
Contact	DAN W MILLER
510(k) history	15 submissions · 12 cleared · 1991-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k970051/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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