

**K091410 DISPOSABLE EMG NEEDLE ELECTRODES, MYOLINE,
MYOLINE 2, MYOBOT,**Aug 11, 2009
90 days to decisionK091410 · Product code: **IKT** · Neurology
Source: <https://www.510kdatabase.net/k091410/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Needle, Diagnostic Electromyograph (IKT)
Date received	May 13, 2009
Decision date	Aug 11, 2009
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Spes Medica Srl
Location	Battipaglia, IT
Contact	ALFREDO SPADAVECCHIA
510(k) history	9 submissions · 9 cleared · 2009-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k091410/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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