

**K092202 ESWALLOW DYSPHAGIA THERAPY KIT, PMI
DYSPHAGIA THERAPY KIT**Feb 10, 2011
568 days to decisionK092202 · Product code: **IPF** · Physical Medicine
Source: <https://www.510kdatabase.net/k092202/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Jul 22, 2009
Decision date	Feb 10, 2011
Days to decision	568 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Eswallow USA
Location	Naples, FL, US
Contact	DANIEL KAMM
510(k) history	2 submissions · 2 cleared · 2011-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k092202/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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