

K093731 VPAP III ST-AMar 22, 2010
109 days to decisionK093731 · Product code: **MNS** · AnesthesiologySource: <https://www.510kdatabase.net/k093731/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Continuous, Non-life-supporting (MNS)
Date received	Dec 3, 2009
Decision date	Mar 22, 2010
Days to decision	109 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Resmed, Ltd.
Location	Poway, CA, US
Contact	DAVID D'CURZ
Website	http://www.resmed.com/
510(k) history	103 submissions · 103 cleared · 1996-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k093731/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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