

**K092085 ISPIRA RESUSCITATION UNIT**Oct 7, 2009  
90 days to decisionK092085 · Product code: **BTL** · Anesthesiology  
Source: <https://www.510kdatabase.net/k092085/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Emergency, Powered (resuscitator) (BTL)
Date received	Jul 9, 2009
Decision date	Oct 7, 2009
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Neoforce Group, Inc.</b>
Location	Doylestown, PA, US
Contact	MONICA FERRANTE
510(k) history	7 submissions · 7 cleared · 2006-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092085/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 24, 2026