

**K924081 LUMISONIC**Nov 9, 1992  
88 days to decisionK924081 · Product code: **CAF** · Anesthesiology  
Source: <https://www.510kdatabase.net/k924081/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Aug 13, 1992
Decision date	Nov 9, 1992
Days to decision	88 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Lumiscope Co., Inc.</b>
Location	Edison, NJ, US
Contact	STEPHEN WHITTENBURG
510(k) history	13 submissions · 13 cleared · 1988-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k924081/> 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 26, 2026