

**K130304 LMA FAMILY OF AIRWAYS**May 30, 2014  
477 days to decisionK130304 · Product code: **CAE** · Anesthesiology  
Source: <https://www.510kdatabase.net/k130304/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Airway, Oropharyngeal, Anesthesiology (CAE)
Date received	Feb 7, 2013
Decision date	May 30, 2014
Days to decision	477 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>The Laryngeal Mask Co., Ltd.</b>
Location	San Diego, CA, US
Contact	PAUL DRYDEN
510(k) history	4 submissions · 4 cleared · 2000-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k130304/> 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