

**K882659 AUGUSTINE INTUBATION GUIDE(TM)**Jul 21, 1988  
23 days to decisionK882659 · Product code: **CCW** · AnesthesiologySource: <https://www.510kdatabase.net/k882659/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laryngoscope, Rigid (CCW)
Date received	Jun 28, 1988
Decision date	Jul 21, 1988
Days to decision	23 days
Third-party review	No

**APPLICANT**

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Company	<b>Augustine Medical, Inc.</b>
Location	Eden Prairie, MN, US
Contact	D AUGUSTINE
510(k) history	25 submissions · 24 cleared · 1987-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882659/>; 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