

**K063159 STERIOX LIQUID CHEMICAL STERILANT SYSTEM**Apr 2, 2007  
167 days to decisionK063159 · Product code: **MED** · General Hospital  
Source: <https://www.510kdatabase.net/k063159/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilant, Medical Devices (MED)
Date received	Oct 17, 2006
Decision date	Apr 2, 2007
Days to decision	167 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Puricore, Inc.</b>
Location	Radnor, PA, US
Contact	HOWARD MANN
510(k) history	10 submissions · 10 cleared · 2006-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k063159/> 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 June 4, 2026