

K013280 STERIOX LIQUID CHEMICAL STERILANT SYSTEM

Sep 18, 2002
 351 days to decision

K013280 · Product code: **MED** · General Hospital
 Source: <https://www.510kdatabase.net/k013280/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilant, Medical Devices (MED)
Date received	Oct 2, 2001
Decision date	Sep 18, 2002
Days to decision	351 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Steriox Technologies, Inc.
Location	Yardley, PA, US
Contact	HOWARD MANN
510(k) history	2 submissions · 2 cleared · 2002-2006

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Device record: <https://www.510kdatabase.net/k013280/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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