FDA 510(k) Premarket Notification

K013280 STERILOX LIQUID CHEMICAL STERILANT SYSTEM

CLEARED

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