FDA 510(k) Premarket Notification

K063159 STERILOX LIQUID CHEMICAL STERILANT SYSTEM

CLEARED

Apr 2, 2007
167 days to decision

K063159 · Product code: **MED** · General Hospital
Source: https://www.510kdatabase.net/k063159/

SUBMISSION DETAILS

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

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