FDA 510(k) Premarket Notification

CLEARED

K090670 MIDMARK M3 ULTRAFAST AUTOMATIC STERILIZER

Jul 7, 2009
116 days to decision

K090670 · Product code: **FLE** · General Hospital
Source: https://www.510kdatabase.net/k090670/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilizer, Steam (FLE)
Date received	Mar 13, 2009
Decision date	Jul 7, 2009
Days to decision	116 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Midmark Corp.**
Location	Versailles, OH, US
Contact	GARY BENNING
510(k) history	31 submissions · 31 cleared · 1981-2013

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k090670/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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