FDA 510(k) Premarket Notification

K140473 DUALGUARD

CLEARED

Apr 10, 2014
44 days to decision

K140473 · Product code: **CCK** · Anesthesiology
Source: https://www.510kdatabase.net/k140473/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Carbon-dioxide, Gaseous-phase (CCK)
Date received	Feb 25, 2014
Decision date	Apr 10, 2014
Days to decision	44 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Flexicare Medical, Ltd.**
Location	Somerset, GB
Contact	CHRISTOPHER WATKINS
510(k) history	7 submissions · 7 cleared · 1995-2015

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k140473/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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