FDA 510(k) Premarket Notification

K142059 FOX MOBILE

CLEARED

Jan 9, 2015
164 days to decision

K142059 · Product code: **CAF** · Anesthesiology
Source: https://www.510kdatabase.net/k142059/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Jul 29, 2014
Decision date	Jan 9, 2015
Days to decision	164 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vectura GmbH**
Location	Gauting, DE
Contact	PAUL DRYDEN
510(k) history	1 submissions · 1 cleared · 2015-2015

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k142059/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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