FDA 510(k) Premarket Notification

K141214 AQUAPAK NEBULIZER ADAPTORS/RESERVOIRS

CLEARED

Dec 18, 2014
220 days to decision

K141214 · Product code: **CAF** · Anesthesiology
Source: https://www.510kdatabase.net/k141214/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	May 12, 2014
Decision date	Dec 18, 2014
Days to decision	220 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Teleflexmedical, Inc.**
Location	Jeffrey, NH, US
Contact	JAMES A COCHIE
510(k) history	64 submissions · 61 cleared · 1985-2024

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k141214/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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