FDA 510(k) Premarket Notification

CLEARED

K140424 RESPIRONICS REUSABLE HEATED TUBING

Nov 14, 2014
268 days to decision

K140424 · Product code: **BZE** · Anesthesiology
Source: https://www.510kdatabase.net/k140424/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Heater, Breathing System W/wo Controller (not Humidifier Or Nebulizer (BZE)
Date received	Feb 19, 2014
Decision date	Nov 14, 2014
Days to decision	268 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Respironics, Inc.**
Location	Monroeville, PA, US
Contact	ALEXANDER L FRIEDMAN
Website	https://www.respironics.com
510(k) history	172 submissions · 168 cleared · 1977-2024

Respironics, Inc. is an American medical supply company owned by Philips. It specializes in products that improve respiratory functions and is based in Monroeville, Pennsylvania. The company maintains a strong FDA 510(k) regulatory record spanning from 1977 to 2024. Respironics has received FDA 510(k) clearances from total submissions. The dominant focus is Anesthesiology devices, which represent approximately 90% of all submissions. The latest clearance in 2024 reflects continued regulatory activity. Recent cleared devices include masks, ventilators, and sleep therapy sy...

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k140424/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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