FDA 510(k) Premarket Notification

K992384 CPAP/PRO CPAP INTERFACE

CLEARED

Dec 1, 1999
138 days to decision

K992384 · Product code: **BZD** · Anesthesiology
Source: https://www.510kdatabase.net/k992384/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Non-continuous (respirator) (BZD)
Date received	Jul 16, 1999
Decision date	Dec 1, 1999
Days to decision	138 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stevenson Industries, Inc.**
Location	Simi Valley, CA, US
Contact	DAVID W SCHLERF
510(k) history	2 submissions · 2 cleared · 1999-2014