FDA 510(k) Premarket Notification

K020944 VORTEX VALVED HOLDING CHAMBER

CLEARED

May 9, 2002
45 days to decision

K020944 · Product code: **CAF** · Anesthesiology
Source: https://www.510kdatabase.net/k020944/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Mar 25, 2002
Decision date	May 9, 2002
Days to decision	45 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Pari Innovative Manufacturers, Inc.**
Location	Midlothian, VA, US
Contact	LAWRENCE WEINSTEIN
510(k) history	10 submissions · 10 cleared · 2002-2007