FDA 510(k) Premarket Notification

K110136 RESPICHAMBER VALVED HOLDING CHAMBER

CLEARED

Apr 18, 2011
90 days to decision

K110136 · Product code: **NVP** · Anesthesiology
Source: https://www.510kdatabase.net/k110136/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Holding Chambers, Direct Patient Interface (NVP)
Date received	Jan 18, 2011
Decision date	Apr 18, 2011
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Trudell Medical Intl.**
Location	London, CA
Contact	DARRYL FISHER
510(k) history	12 submissions · 12 cleared · 2002-2013