FDA 510(k) Premarket Notification

K142096 T-PIECE RESUSCITATOR

CLEARED

Oct 17, 2014
77 days to decision

K142096 · Product code: **BTL** · Anesthesiology
Source: https://www.510kdatabase.net/k142096/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Emergency, Powered (resuscitator) (BTL)
Date received	Aug 1, 2014
Decision date	Oct 17, 2014
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Mercury Medical**
Location	Clearwater, FL, US
Contact	PAUL DRYDEN
510(k) history	15 submissions · 15 cleared · 1994-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k142096/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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