FDA 510(k) Premarket Notification

K141940 CONCHASMART COLUMN

CLEARED

Nov 24, 2014
130 days to decision

K141940 · Product code: **BTT** · Anesthesiology
Source: https://www.510kdatabase.net/k141940/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Humidifier, Respiratory Gas, (direct Patient Interface) (BTT)
Date received	Jul 17, 2014
Decision date	Nov 24, 2014
Days to decision	130 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Teleflex, Inc.**
Location	Jeffrey, NH, US
Contact	Amanda Webb
510(k) history	10 submissions · 10 cleared · 1986-2016