

K141940 CONCHASMART COLUMNNov 24, 2014
130 days to decisionK141940 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k141940/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 24, 2026