FDA 510(k) Premarket Notification

K813075 TRITEC DOC TM

CLEARED

Dec 9, 1981
37 days to decision

K813075 · Product code: **BZD** · Anesthesiology
Source: https://www.510kdatabase.net/k813075/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Non-continuous (respirator) (BZD)
Date received	Nov 2, 1981
Decision date	Dec 9, 1981
Days to decision	37 days
Third-party review	No

APPLICANT

Company	**Tri-Tech, Inc.**
Location	Mchenry, IL, US
510(k) history	11 submissions · 9 cleared · 1981-1997