

K813075 TRITEC DOC TMDec 9, 1981
37 days to decisionK813075 · 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

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Device record: <https://www.510kdatabase.net/k813075/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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