FDA 510(k) Premarket Notification

K810219 PREFILLED HUMIDIFER

CLEARED

Mar 4, 1981
36 days to decision

K810219 · Product code: **BTT** · Anesthesiology
Source: https://www.510kdatabase.net/k810219/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Humidifier, Respiratory Gas, (direct Patient Interface) (BTT)
Date received	Jan 27, 1981
Decision date	Mar 4, 1981
Days to decision	36 days
Third-party review	No

APPLICANT

Company	**Corpak Co.**
Location	Mchenry, IL, US
510(k) history	21 submissions · 20 cleared · 1981-1996