

K810219 PREFILLED HUMIDIFERMar 4, 1981
36 days to decisionK810219 · Product code: **BTT** · AnesthesiologySource: <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

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Device record: <https://www.510kdatabase.net/k810219/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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