

K822816 DUAL SERVO HEATED RESPIRATORY HUMIDIFIEROct 22, 1982
32 days to decisionK822816 · Product code: **BTT** · AnesthesiologySource: <https://www.510kdatabase.net/k822816/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Humidifier, Respiratory Gas, (direct Patient Interface) (BTT)
Date received	Sep 20, 1982
Decision date	Oct 22, 1982
Days to decision	32 days
Third-party review	No

APPLICANT

Company	Fisher & Paykel Allied Products , Ltd.
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1982-1988

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k822816/> 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 27, 2026