

**K020899 PARAPAC 200 MEDIC, MODEL P200, & PARAPAC
200D MEDIC, MODEL P200D**Dec 13, 2002
268 days to decisionK020899 · Product code: **BTL** · Anesthesiology
Source: <https://www.510kdatabase.net/k020899/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Emergency, Powered (resuscitator) (BTL)
Date received	Mar 20, 2002
Decision date	Dec 13, 2002
Days to decision	268 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Pneupac , Ltd.
Location	Walker, MI, US
Contact	DONALD ALEXANDER
510(k) history	19 submissions · 19 cleared · 1980-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k020899/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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