FDA 510(k) Premarket Notification

K820220 VENTILATOR MODELS

CLEARED

Feb 18, 1982
22 days to decision

K820220 · Product code: **CBK** · Anesthesiology
Source: https://www.510kdatabase.net/k820220/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Continuous, Facility Use (CBK)
Date received	Jan 27, 1982
Decision date	Feb 18, 1982
Days to decision	22 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k820220/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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