

K792452 PNEUPAC RESUSCITATOR INSTANT ACTION SETFeb 5, 1980
64 days to decisionK792452 · Product code: **DRM** · CardiovascularSource: <https://www.510kdatabase.net/k792452/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Compressor, Cardiac, External (DRM)
Date received	Dec 3, 1979
Decision date	Feb 5, 1980
Days to decision	64 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.netDevice record: <https://www.510kdatabase.net/k792452/> 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 25, 2026