

**K792451 PNEUPAC RESUSCITATOR INSTANT ACTION SET**Feb 5, 1980  
64 days to decisionK792451 · Product code: **DRM** · CardiovascularSource: <https://www.510kdatabase.net/k792451/>**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	<b>Pneupac , Ltd.</b>
Location	Walker, MI, US
510(k) history	19 submissions · 19 cleared · 1980-2004

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k792451/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 25, 2026