FDA 510(k) Premarket Notification

K792452 PNEUPAC RESUSCITATOR INSTANT ACTION SET

CLEARED

Feb 5, 1980
64 days to decision

K792452 · Product code: **DRM** · Cardiovascular
Source: 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