

K072372 PHLEBOPRESS MODEL 601A COMPRESSION THERAPY DEVICENov 21, 2007
90 days to decisionK072372 · Product code: **JOW** · Cardiovascular
Source: <https://www.510kdatabase.net/k072372/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sleeve, Limb, Compressible (JOW)
Date received	Aug 23, 2007
Decision date	Nov 21, 2007
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Mego Afek
Location	Walker, MI, US
Contact	AHAVA STEIN
510(k) history	5 submissions · 5 cleared · 1981-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k072372/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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