FDA 510(k) Premarket Notification

K072372 PHLEBOPRESS MODEL 601A COMPRESSION THERAPY DEVICE

CLEARED

Nov 21, 2007
90 days to decision

K072372 · 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