FDA 510(k) Premarket Notification

K122281 PADNET 2.0

CLEARED

Aug 29, 2012
30 days to decision

K122281 · Product code: **JOM** · Cardiovascular
Source: https://www.510kdatabase.net/k122281/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Photoelectric, Pneumatic Or Hydraulic (JOM)
Date received	Jul 30, 2012
Decision date	Aug 29, 2012
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Biomedix, Inc.**
Location	Spencer, IN, US
Contact	GREG HOCKING
510(k) history	11 submissions · 11 cleared · 1993-2014