

K122281 PADNET 2.0Aug 29, 2012
30 days to decisionK122281 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k122281/> Data sourced from FDA 510(k) public records (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. - Generated May 20, 2026