FDA 510(k) Premarket Notification

K073148 IDRIVES, IDRIVEC, IDRIVEF WITH VASCULAR INDICATIONS

CLEARED

Dec 7, 2007
29 days to decision

K073148 · Product code: **GDW** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k073148/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Staple, Implantable (GDW)
Date received	Nov 8, 2007
Decision date	Dec 7, 2007
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Power Medical Interventions, Inc.**
Location	New Hope, PA, US
Contact	BARBARA J WHITMAN
510(k) history	22 submissions · 22 cleared · 2001-2009

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k073148/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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