

K092382 NEXUS PRESSURE RATED EXTENSION SETSApr 26, 2010
264 days to decisionK092382 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k092382/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 5, 2009
Decision date	Apr 26, 2010
Days to decision	264 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Nexus Medical, LLC
Location	Raleigh, NC, US
Contact	SUSAN CURRY
510(k) history	16 submissions · 16 cleared · 2003-2015

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k092382/> 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 June 5, 2026