

**K081515 NEOMED DUAL LUMEN UMBILICAL CATHETER,
MODELS S2UVC4.0, S2UVC5.0**

Jul 21, 2008
52 days to decision

K081515 · Product code: **FOS** · General Hospital
Source: <https://www.510kdatabase.net/k081515/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Umbilical Artery (FOS)
Date received	May 30, 2008
Decision date	Jul 21, 2008
Days to decision	52 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Neomed, Inc.
Location	Swanee, GA, US
Contact	PENNY NORTHCUTT
510(k) history	13 submissions · 13 cleared · 2007-2019

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k081515/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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