

K091213 FOOTPRINT MEDICAL UMBILICAL CATHETER, DUAL LUMENSep 25, 2009
151 days to decisionK091213 · Product code: **FOS** · General Hospital
Source: <https://www.510kdatabase.net/k091213/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Umbilical Artery (FOS)
Date received	Apr 27, 2009
Decision date	Sep 25, 2009
Days to decision	151 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Footprint Medical
Location	San Antonio, TX, US
Contact	CLYDE BAKER
510(k) history	6 submissions · 6 cleared · 2009-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k091213/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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