

**K122263 14.5-16.0 FRENCH CUFFED CATHETER REMOVAL
DEVICE, SYMMETRY SURGICAL SINGEL USE**Sep 28, 2012
63 days to decisionK122263 · Product code: **ODY** · General Hospital
Source: <https://www.510kdatabase.net/k122263/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tunneled Catheter Remover (ODY)
Date received	Jul 27, 2012
Decision date	Sep 28, 2012
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Symmetry Medical
Location	New Bedford, MA, US
Contact	HANNAH FOLEY
510(k) history	2 submissions · 2 cleared · 2012-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k122263/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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