

K042491 BD ONECATH PERIPHERALLY INSERTED CENTRAL CATHETERNov 22, 2004
69 days to decisionK042491 · Product code: **LJS** · General Hospital
Source: <https://www.510kdatabase.net/k042491/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Sep 14, 2004
Decision date	Nov 22, 2004
Days to decision	69 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Becton, Dickinson & CO
Location	Franklin Lakes, NJ, US
Contact	LESLIE WOOD
510(k) history	190 submissions · 190 cleared · 2001-2016

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k042491/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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