

K051194 BD ONECATH PERIPHERALLY INSERTED CENTRAL CATHETER, MODELS 384647, 384667, 384687, 384648, 384668, 384688, 384649, 384669

Jul 25, 2005
76 days to decision

K051194 · Product code: **LJS** · General Hospital
Source: <https://www.510kdatabase.net/k051194/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	May 10, 2005
Decision date	Jul 25, 2005
Days to decision	76 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

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Device record: <https://www.510kdatabase.net/k051194/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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