

**K882448 MULTILUMEN CENTRAL VENOUS CATH. W/VITACUFF
DEVICE**Aug 9, 1988
56 days to decisionK882448 · Product code: **LJS** · General Hospital
Source: <https://www.510kdatabase.net/k882448/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Jun 14, 1988
Decision date	Aug 9, 1988
Days to decision	56 days
Third-party review	No

APPLICANT

Company	Bd Becton Dickinson Vacutainer Systems Preanalytic
Location	Washington, DC, US
Contact	CHARLES J WELLE
510(k) history	632 submissions · 625 cleared · 1976-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k882448/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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