

**K893126 DESERET(R) INTRODUCER SET WITH VITACUFF(R)
DEVICE**Aug 10, 1989
107 days to decisionK893126 · Product code: **LJS** · General Hospital
Source: <https://www.510kdatabase.net/k893126/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - SD
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Apr 25, 1989
Decision date	Aug 10, 1989
Days to decision	107 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/k893126/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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