FDA 510(k) Premarket Notification

K893126 DESERET(R) INTRODUCER SET WITH VITACUFF(R) DEVICE

CLEARED

Aug 10, 1989
107 days to decision

K893126 · 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