FDA 510(k) Premarket Notification

K882448 MULTILUMEN CENTRAL VENOUS CATH. W/VITACUFF DEVICE

CLEARED

Aug 9, 1988
56 days to decision

K882448 · 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