FDA 510(k) Premarket Notification

K880776 7 FR. TRIPLE LUMEN CENTRAL VEIN CATHETERI. SYSTEM

CLEARED

May 2, 1988
66 days to decision

K880776 · Product code: **DQR** · Cardiovascular
Source: https://www.510kdatabase.net/k880776/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Catheter (DQR)
Date received	Feb 26, 1988
Decision date	May 2, 1988
Days to decision	66 days
Third-party review	No

APPLICANT

Company	**Medsurg Industries, Inc.**
Location	Mchenry, IL, US
Contact	TOM BONNER
510(k) history	41 submissions · 26 cleared · 1982-1994