FDA 510(k) Premarket Notification

K911519 CATHCAP, CAT. NO. CC100 SERIES

CLEARED

May 22, 1991
48 days to decision

K911519 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k911519/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Apr 4, 1991
Decision date	May 22, 1991
Days to decision	48 days
Third-party review	No

APPLICANT

Company	**Gish Biomedical, Inc.**
Location	Mchenry, IL, US
Contact	DEBI KRIDNER
510(k) history	75 submissions · 75 cleared · 1983-2009