FDA 510(k) Premarket Notification

K895942 GESCO DUAL LUMEN PER-Q-CATH-DL

CLEARED

Jul 2, 1990
264 days to decision

K895942 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k895942/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Oct 11, 1989
Decision date	Jul 2, 1990
Days to decision	264 days
Third-party review	No

APPLICANT

Company	**Gesco Intl., Inc.**
Location	Walker, MI, US
Contact	GEORGE E SINKO
510(k) history	26 submissions · 22 cleared · 1978-1995