FDA 510(k) Premarket Notification

K831845 PER-Q-CATH TRAY

CLEARED

Aug 24, 1983
77 days to decision

K831845 · Product code: **LJS** · General Hospital
Source: https://www.510kdatabase.net/k831845/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Jun 8, 1983
Decision date	Aug 24, 1983
Days to decision	77 days
Third-party review	No

APPLICANT

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