FDA 510(k) Premarket Notification

K901419 DUAL PORT

CLEARED

Sep 5, 1990
163 days to decision

K901419 · Product code: **LJT** · General Hospital
Source: https://www.510kdatabase.net/k901419/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Mar 26, 1990
Decision date	Sep 5, 1990
Days to decision	163 days
Third-party review	No

APPLICANT

Company	**Infusaid Corp.**
Location	Walker, MI, US
Contact	NANCY C HALL
510(k) history	8 submissions · 8 cleared · 1983-1991