FDA 510(k) Premarket Notification

K882230 A PORT IMPLANT VASCULAR ACCESS SYSTEM - DETACHED

CLEARED

Jul 29, 1988
63 days to decision

K882230 · Product code: **LJT** · General Hospital
Source: https://www.510kdatabase.net/k882230/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	May 27, 1988
Decision date	Jul 29, 1988
Days to decision	63 days
Third-party review	No

APPLICANT

Company	**Therex Corp.**
Location	Walpole, MA, US
Contact	BRADLEY J ENEGREN
510(k) history	26 submissions · 23 cleared · 1988-1995