FDA 510(k) Premarket Notification

K830000 VASCULAR-ACCESS-PORT

CLEARED

Mar 1, 1983
57 days to decision

K830000 · Product code: **LJT** · General Hospital
Source: https://www.510kdatabase.net/k830000/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Jan 3, 1983
Decision date	Mar 1, 1983
Days to decision	57 days
Third-party review	No

APPLICANT

Company	**Norfolk Medical Products, Inc.**
Location	Walker, MI, US
510(k) history	20 submissions · 20 cleared · 1983-2020