FDA 510(k) Premarket Notification

K951814 TRIUMPH-1 PORT

CLEARED

Aug 29, 1995
133 days to decision

K951814 · Product code: **LJT** · General Hospital
Source: https://www.510kdatabase.net/k951814/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Apr 18, 1995
Decision date	Aug 29, 1995
Days to decision	133 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Horizon Medical Products, Inc.**
Location	Atlanta, GA, US
Contact	RODDY J. H CLARK
510(k) history	16 submissions · 11 cleared · 1994-2003

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k951814/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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