FDA 510(k) Premarket Notification

K934640 HMP DUAL POLYSULFONE PORT

CLEARED

Feb 25, 1994
150 days to decision

K934640 · Product code: **LJT** · General Hospital
Source: https://www.510kdatabase.net/k934640/

SUBMISSION DETAILS

Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Sep 28, 1993
Decision date	Feb 25, 1994
Days to decision	150 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