FDA 510(k) Premarket Notification

K960350 THERAPORT VASCULAR ACCESS SYSTEM

CLEARED

Apr 15, 1996
81 days to decision

K960350 · Product code: **LJT** · General Hospital
Source: https://www.510kdatabase.net/k960350/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Jan 25, 1996
Decision date	Apr 15, 1996
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary
Other names	MODEL 1001, 1002, 1601, 1602

APPLICANT

Company	**Biocontrol Technology, Inc.**
Location	Indiana, PA, US
Contact	PATRICK J COOPER
510(k) history	2 submissions · 2 cleared · 1996-1998