

**K960350 THERAPORT VASCULAR ACCESS SYSTEM**Apr 15, 1996  
81 days to decision

K960350 · Product code: LJT · General Hospital

Source: <https://www.510kdatabase.net/k960350/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960350/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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