

**K013635 TRELIS INFUSION SYSTEM (10CM INFUSION LENGTH)**

Feb 11, 2002  
98 days to decision

K013635 · Product code: **QEY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k013635/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mechanical Thrombolysis Catheter (QEY)
Date received	Nov 5, 2001
Decision date	Feb 11, 2002
Days to decision	98 days
Third-party review	No
Summary / Statement	Summary
Other names	TRELIS INFUSION SYSTEM (20CM INFUSION LENGTH)

**APPLICANT**

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Company	<b>Bacchus Vascular, Inc.</b>
Location	Santa Clara, CA, US
Contact	GREG MATHISON
510(k) history	10 submissions · 10 cleared · 2001-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k013635/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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