

**K963957 TRINITY DVT SYSTEM**Nov 4, 1997  
398 days to decisionK963957 · Product code: **JOW** · Cardiovascular  
Source: <https://www.510kdatabase.net/k963957/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sleeve, Limb, Compressible (JOW)
Date received	Oct 2, 1996
Decision date	Nov 4, 1997
Days to decision	398 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Delphi Consulting Group</b>
Location	Houston, TX, US
Contact	J. HARVEY KNAUSS
510(k) history	6 submissions · 6 cleared · 1996-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k963957/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 4, 2026