

**K930286 DRI-VENDT/PROVENT/ACCUVENT ARTERIAL BLOOD  
GAS KITS**Jun 22, 1993  
152 days to decisionK930286 · Product code: **CBT** · Anesthesiology  
Source: <https://www.510kdatabase.net/k930286/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - KD
Submission type	Traditional
Device classification	Arterial Blood Sampling Kit (CBT)
Date received	Jan 21, 1993
Decision date	Jun 22, 1993
Days to decision	152 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Concord/Portex</b>
Location	Keene, NH, US
Contact	ROBERT WHEELER
510(k) history	23 submissions · 20 cleared · 1989-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k930286/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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