

**K033323 D 903 AVANT ADULT HOLLOW FIBER OXYGENATOR  
AND D 903 AVANT 2 PH.I.S.I.O. ADULT HOLLOW FIBER  
OXYGENATOR WITH COATING**Jan 13, 2004  
90 days to decisionK033323 · Product code: **DTZ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k033323/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Oxygenator, Cardiopulmonary Bypass (DTZ)
Date received	Oct 15, 2003
Decision date	Jan 13, 2004
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Dideco S.P.A.</b>
Location	Waltham, MA, US
Contact	BARRY SALL
510(k) history	22 submissions · 22 cleared · 1997-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k033323/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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