

**K032040 IDEAL MIMESYS SYSTEM WITH INTEGRATED
VENOUS AIR REMOVAL, CENTRIFUGAL BLOOD PUMP, PUMP
BRACKET, ADULT MEMBRANE OXYGENATOR**Jul 9, 2003
8 days to decisionK032040 · Product code: **KFM** · Cardiovascular
Source: <https://www.510kdatabase.net/k032040/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Pump, Blood, Cardiopulmonary Bypass, Non-roller Type (KFM)
Date received	Jul 1, 2003
Decision date	Jul 9, 2003
Days to decision	8 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k032040/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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