

**K953835 LILLIPUT 1 NEW BORN HOLLOW FIBER
OXYGENATOR**Mar 15, 1996
212 days to decisionK953835 · Product code: **DTZ** · Cardiovascular
Source: <https://www.510kdatabase.net/k953835/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oxygenator, Cardiopulmonary Bypass (DTZ)
Date received	Aug 16, 1995
Decision date	Mar 15, 1996
Days to decision	212 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Dideco S.R.L.
Location	Waltham, MA, US
Contact	BARRY SALL
510(k) history	6 submissions · 6 cleared · 1996-2005

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k953835/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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