

K951369 EDMNov 7, 1995
225 days to decisionK951369 · Product code: **DPW** · CardiovascularSource: <https://www.510kdatabase.net/k951369/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	Mar 27, 1995
Decision date	Nov 7, 1995
Days to decision	225 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Deltex Medical
Location	Rockville, MD, US
Contact	GLENN A RAHMOELLER
510(k) history	3 submissions · 3 cleared · 1995-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k951369/> Data sourced from FDA 510(k) public records (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. - Generated June 28, 2026