

K982385 POROUS MEDIA DPB20 SERIES DISPOSABLE PRE-BYPASS FILTERFeb 1, 1999
207 days to decisionK982385 · Product code: **KRJ** · Cardiovascular
Source: <https://www.510kdatabase.net/k982385/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Prebypass, Cardiopulmonary Bypass (KRJ)
Date received	Jul 9, 1998
Decision date	Feb 1, 1999
Days to decision	207 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Porous Media Corp.
Location	St. Paul, MN, US
Contact	ROCHELLE M MICKSCHL
510(k) history	9 submissions · 9 cleared · 1994-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k982385/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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