

K913728 DENVER(R) PERITONEO-SUBCLAVIAN SHUNTFeb 18, 1994
913 days to decisionK913728 · Product code: **KPM** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k913728/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shunt, Peritoneal (KPM)
Date received	Aug 20, 1991
Decision date	Feb 18, 1994
Days to decision	913 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Denver Biomedicals, Inc.
Location	Englewood, CO, US
Contact	SHIRLEY K FREEMAN
510(k) history	10 submissions · 10 cleared · 1981-2005

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k913728/> 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 May 25, 2026