FDA 510(k) Premarket Notification

K913728 DENVER(R) PERITONEO-SUBCLAVIAN SHUNT

CLEARED

Feb 18, 1994
913 days to decision

K913728 · Product code: **KPM** · Gastroenterology & Urology
Source: 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