FDA 510(k) Premarket Notification

K811183 DENVER PERITONEO-VENOUS SHUNT

CLEARED

Jul 1, 1981
63 days to decision

K811183 · Product code: **KPM** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k811183/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shunt, Peritoneal (KPM)
Date received	Apr 29, 1981
Decision date	Jul 1, 1981
Days to decision	63 days
Third-party review	No

APPLICANT

Company	**Denver Biomedicals, Inc.**
Location	Englewood, CO, US
Contact	JOHN B NEWKIRK
510(k) history	10 submissions · 10 cleared · 1981-2005