FDA 510(k) Premarket Notification

K880401 DENVER PERITONEO-VENOUS SHUNT

CLEARED

May 31, 1988
123 days to decision

K880401 · Product code: **KPM** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k880401/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shunt, Peritoneal (KPM)
Date received	Jan 29, 1988
Decision date	May 31, 1988
Days to decision	123 days
Third-party review	No

APPLICANT

Company	**Codman & Shurtleff, Inc.**
Location	Mchenry, IL, US
Contact	THOMAS J COURAGE
510(k) history	152 submissions · 151 cleared · 1976-2020