FDA 510(k) Premarket Notification

K880570 DENVER PLEURO-PERITONEAL SHUNT

CLEARED

Jun 20, 1988
131 days to decision

K880570 · Product code: **KPM** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k880570/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shunt, Peritoneal (KPM)
Date received	Feb 10, 1988
Decision date	Jun 20, 1988
Days to decision	131 days
Third-party review	No

APPLICANT

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