FDA 510(k) Premarket Notification

K900884 P.R.G. PERCUTANEOUS REPLACEMENT GASTROSTOMY

CLEARED

Jul 2, 1990
126 days to decision

K900884 · Product code: **KGC** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k900884/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SD
Submission type	Traditional
Device classification	Tube, Gastro-enterostomy (KGC)
Date received	Feb 26, 1990
Decision date	Jul 2, 1990
Days to decision	126 days
Third-party review	No

APPLICANT

Company	**Applied Medical Technologies**
Location	Mchenry, IL, US
Contact	PAT GILPIN
510(k) history	23 submissions · 20 cleared · 1981-1998