FDA 510(k) Premarket Notification

K800186 LOGO-NU-WAY

CLEARED

Feb 26, 1980
28 days to decision

K800186 · Product code: **KNX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k800186/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Urine, (and Accessories) For Indwelling Catheter (KNX)
Date received	Jan 29, 1980
Decision date	Feb 26, 1980
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Gary Giancarlo**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980