FDA 510(k) Premarket Notification

K810645 SELECT-A-FLOW COLLECTOR

CLEARED

May 5, 1981
56 days to decision

K810645 · Product code: **KNX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k810645/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Urine, (and Accessories) For Indwelling Catheter (KNX)
Date received	Mar 10, 1981
Decision date	May 5, 1981
Days to decision	56 days
Third-party review	No

APPLICANT

Company	**Wenmar Intl. Corp.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1981-1981