FDA 510(k) Premarket Notification

K781422 KWIK-FLO URINAL SYSTEM

CLEARED

Sep 7, 1978
21 days to decision

K781422 · Product code: **KNX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k781422/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Urine, (and Accessories) For Indwelling Catheter (KNX)
Date received	Aug 17, 1978
Decision date	Sep 7, 1978
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**The Grace Mfg. Co.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978