FDA 510(k) Premarket Notification

K791196 URIDROP

CLEARED

Jul 30, 1979
34 days to decision

K791196 · Product code: **KNX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k791196/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Urine, (and Accessories) For Indwelling Catheter (KNX)
Date received	Jun 26, 1979
Decision date	Jul 30, 1979
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Ab Medett Produkter**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1979-1979