FDA 510(k) Premarket Notification

K862251 MODIFIED BIO-FLOW MODEL 200 URINE FLOW DEVICE

CLEARED

Sep 4, 1986
84 days to decision

K862251 · Product code: **KNX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k862251/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Urine, (and Accessories) For Indwelling Catheter (KNX)
Date received	Jun 12, 1986
Decision date	Sep 4, 1986
Days to decision	84 days
Third-party review	No

APPLICANT

Company	**Bio-Flow, Inc.**
Location	Costa Mesa, CA, US
Contact	THOMAS E HYANS
510(k) history	2 submissions · 2 cleared · 1985-1986