FDA 510(k) Premarket Notification

K884365 MULTIFLOW(TM) 60 KIT

CLEARED

Dec 8, 1988
51 days to decision

K884365 · Product code: **KDI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k884365/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Oct 18, 1988
Decision date	Dec 8, 1988
Days to decision	51 days
Third-party review	No

APPLICANT

Company	**Hospal Medical Corp.**
Location	Mchenry, IL, US
Contact	CATHERINA MADORMO
510(k) history	55 submissions · 55 cleared · 1977-1989