FDA 510(k) Premarket Notification

CLEARED

K883307 CARDIOFLOW 100 HEMOCONCENTRATOR/PREPARATION KIT

Oct 12, 1988
69 days to decision

K883307 · Product code: **KDI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k883307/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Aug 4, 1988
Decision date	Oct 12, 1988
Days to decision	69 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