FDA 510(k) Premarket Notification

K852885 FILTRAL HEMODIALYZER

CLEARED

Sep 6, 1985
59 days to decision

K852885 · Product code: **KDI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k852885/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Jul 9, 1985
Decision date	Sep 6, 1985
Days to decision	59 days
Third-party review	No

APPLICANT

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