FDA 510(k) Premarket Notification

K872456 HEMOCOR PLUS HEMOCONCENTRATOR

CLEARED

Sep 15, 1987
84 days to decision

K872456 · Product code: **KDI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k872456/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Jun 23, 1987
Decision date	Sep 15, 1987
Days to decision	84 days
Third-party review	No

APPLICANT

Company	**Minntech Corp.**
Location	Minneapolis, MN, US
Contact	LEROY J FISCHBACH
510(k) history	33 submissions · 33 cleared · 1987-2012