FDA 510(k) Premarket Notification

K980859 HEMOCOR HPH MINI HEMOCONCENTRATOR

CLEARED

May 12, 1998
68 days to decision

K980859 · Product code: **KDI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k980859/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Mar 5, 1998
Decision date	May 12, 1998
Days to decision	68 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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