FDA 510(k) Premarket Notification

K923139 HEMOCOR HP HEMOCONCENTRATOR SERIES

CLEARED

Feb 18, 1994
599 days to decision

K923139 · Product code: **KDI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k923139/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Jun 29, 1992
Decision date	Feb 18, 1994
Days to decision	599 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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