FDA 510(k) Premarket Notification

K071298 HF JUNIOR HEMOFILTER

CLEARED

Aug 8, 2007
91 days to decision

K071298 · Product code: **KDI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k071298/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	May 9, 2007
Decision date	Aug 8, 2007
Days to decision	91 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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