

K071298 HF JUNIOR HEMOFILTERAug 8, 2007
91 days to decisionK071298 · Product code: **KDI** · Gastroenterology & UrologySource: <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

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