FDA 510(k) Premarket Notification

K830595 MONITRAL

CLEARED

Apr 28, 1983
63 days to decision

K830595 · Product code: **KDI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k830595/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Feb 24, 1983
Decision date	Apr 28, 1983
Days to decision	63 days
Third-party review	No

APPLICANT

Company	**Hospal Medical Corp.**
Location	Mchenry, IL, US
510(k) history	55 submissions · 55 cleared · 1977-1989