FDA 510(k) Premarket Notification

K823589 H 12 07

CLEARED

Dec 22, 1982
16 days to decision

K823589 · Product code: **KDI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k823589/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Dec 6, 1982
Decision date	Dec 22, 1982
Days to decision	16 days
Third-party review	No

APPLICANT

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