FDA 510(k) Premarket Notification

K890315 CAPILLARY FLOW DIALYZERS MODELS CT110G & CT190G

CLEARED

Feb 8, 1989
16 days to decision

K890315 · Product code: **KDI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k890315/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Jan 23, 1989
Decision date	Feb 8, 1989
Days to decision	16 days
Third-party review	No

APPLICANT

Company	**Baxter Healthcare Corp**
Location	Mchenry, IL, US
Contact	ROBERT L WILKINSON
510(k) history	505 submissions · 496 cleared · 1977-2019