FDA 510(k) Premarket Notification

K821834 MODEL 100 BICARBONATE CONDUCTIVITY

CLEARED

Jul 2, 1982
10 days to decision

K821834 · Product code: **FKQ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k821834/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Dialysate Delivery, Central Multiple Patient (FKQ)
Date received	Jun 22, 1982
Decision date	Jul 2, 1982
Days to decision	10 days
Third-party review	No

APPLICANT

Company	**Western Laboratories Corp.**
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1978-1984