FDA 510(k) Premarket Notification

K800666 QUIK-TEMP

CLEARED

May 8, 1980
43 days to decision

K800666 · Product code: **KPP** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k800666/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Peritoneal Dialysate Filter (KPP)
Date received	Mar 26, 1980
Decision date	May 8, 1980
Days to decision	43 days
Third-party review	No

APPLICANT

Company	**Liquid Crystal Products, Inc.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1980-1982