

K800666 QUIK-TEMPMay 8, 1980
43 days to decisionK800666 · Product code: **KPP** · Gastroenterology & UrologySource: <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

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Device record: <https://www.510kdatabase.net/k800666/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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