

**K812615 TM-CAPD**Jan 5, 1982  
112 days to decisionK812615 · Product code: **KPP** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k812615/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Peritoneal Dialysate Filter (KPP)
Date received	Sep 15, 1981
Decision date	Jan 5, 1982
Days to decision	112 days
Third-party review	No

**APPLICANT**

---

Company	<b>Tri-Med, Inc.</b>
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1977-2004

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k812615/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 20, 2026