

**K811981 ISOLATER THP**Aug 4, 1981  
22 days to decisionK811981 · Product code: **KOC** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k811981/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Blood Circuit, Hemodialysis (KOC)
Date received	Jul 13, 1981
Decision date	Aug 4, 1981
Days to decision	22 days
Third-party review	No

**APPLICANT**

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

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Device record: <https://www.510kdatabase.net/k811981/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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