

**K830257 NON-STERILE PERITONEAL ACCESS CATHET-**Feb 24, 1983  
30 days to decisionK830257 · Product code: **FKX** · Microbiology  
Source: <https://www.510kdatabase.net/k830257/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Peritoneal, Automatic Delivery (FKX)
Date received	Jan 25, 1983
Decision date	Feb 24, 1983
Days to decision	30 days
Third-party review	No

**APPLICANT**

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Company	<b>Evermed, Inc.</b>
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1983-1984

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Device record: <https://www.510kdatabase.net/k830257/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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