

**K780470 H.P. MEDICAL GRADE SILICONE TUBING**Apr 27, 1978  
36 days to decisionK780470 · Product code: **KDL** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k780470/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Perfusion, Kidney, Disposable (KDL)
Date received	Mar 22, 1978
Decision date	Apr 27, 1978
Days to decision	36 days
Third-party review	No

**APPLICANT**

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Company	<b>Dow Corning Corp. Healthcare Industries Materials</b>
Location	Mchenry, IL, US
510(k) history	31 submissions · 31 cleared · 1976-1985

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

Device record: <https://www.510kdatabase.net/k780470/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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