

**K761007 ARTIFICIAL KIDNEY, C-DAK CORDIS DOW**Jan 12, 1977  
65 days to decisionK761007 · Product code: **FJI** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k761007/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, Capillary, Hollow Fiber (FJI)
Date received	Nov 8, 1976
Decision date	Jan 12, 1977
Days to decision	65 days
Third-party review	No

**APPLICANT**

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Company	<b>Cordis Corp.</b>
Location	Mchenry, IL, US
Website	<a href="https://cordis.com">https://cordis.com</a>
510(k) history	315 submissions · 281 cleared · 1976-2014

Cordis Corp. is a medical device manufacturer based in McHenry, US. The company specializes in interventional cardiovascular and gastroenterology devices. Cordis has a substantial FDA 510(k) regulatory history spanning from 1976 to 2014. The company received FDA 510(k) clearances from total submissions. Its portfolio focuses primarily on cardiovascular devices and gastroenterology stent systems, including percutaneous transluminal angioplasty catheters, emboli capture guidewires, and self-expanding biliary stent systems. Notable cleared products include the FLEXSTENT Bili...

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