

**K890315 CAPILLARY FLOW DIALYZERS MODELS CT110G & CT190G**Feb 8, 1989  
16 days to decisionK890315 · Product code: **KDI** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k890315/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Jan 23, 1989
Decision date	Feb 8, 1989
Days to decision	16 days
Third-party review	No

**APPLICANT**

---

Company	<b>Baxter Healthcare Corp</b>
Location	Mchenry, IL, US
Contact	ROBERT L WILKINSON
510(k) history	505 submissions · 496 cleared · 1977-2019

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890315/> Data sourced from FDA 510(k) public records (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. - Generated May 17, 2026