

**K862251 MODIFIED BIO-FLOW MODEL 200 URINE FLOW DEVICE**Sep 4, 1986  
84 days to decisionK862251 · Product code: **KNX** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k862251/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Urine, (and Accessories) For Indwelling Catheter (KNX)
Date received	Jun 12, 1986
Decision date	Sep 4, 1986
Days to decision	84 days
Third-party review	No

**APPLICANT**

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Company	<b>Bio-Flow, Inc.</b>
Location	Costa Mesa, CA, US
Contact	THOMAS E HYANS
510(k) history	2 submissions · 2 cleared · 1985-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k862251/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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