

K821829 UROGARDJul 30, 1982
38 days to decisionK821829 · Product code: **KNX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k821829/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Urine, (and Accessories) For Indwelling Catheter (KNX)
Date received	Jun 22, 1982
Decision date	Jul 30, 1982
Days to decision	38 days
Third-party review	No

APPLICANT

Company	Buckman Consultants
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1982-1983

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Device record: <https://www.510kdatabase.net/k821829/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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