

K791196 URIDROPJul 30, 1979
34 days to decisionK791196 · Product code: **KNX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k791196/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Urine, (and Accessories) For Indwelling Catheter (KNX)
Date received	Jun 26, 1979
Decision date	Jul 30, 1979
Days to decision	34 days
Third-party review	No

APPLICANT

Company	Ab Medett Produkter
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1979-1979

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Device record: <https://www.510kdatabase.net/k791196/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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