

K800505 URETHRAL CATHETERApr 8, 1980
35 days to decisionK800505 · Product code: **KOD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k800505/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urological (KOD)
Date received	Mar 4, 1980
Decision date	Apr 8, 1980
Days to decision	35 days
Third-party review	No

APPLICANT

Company	Medi-Craft , Ltd.
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1980-1986

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Device record: <https://www.510kdatabase.net/k800505/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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