

K791888 CATHETER CARE KIT REORDER 7940&7941Oct 17, 1979
36 days to decisionK791888 · Product code: **EFQ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k791888/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Gauze/sponge, Internal (EFQ)
Date received	Sep 11, 1979
Decision date	Oct 17, 1979
Days to decision	36 days
Third-party review	No

APPLICANT

Company	Ascep, Inc.
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1979-1983

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Device record: <https://www.510kdatabase.net/k791888/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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