

K801632 THE BELLCO FISTULA NEEDLESAug 4, 1980
20 days to decisionK801632 · Product code: **FIE** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k801632/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	Jul 15, 1980
Decision date	Aug 4, 1980
Days to decision	20 days
Third-party review	No

APPLICANT

Company	Bellco Artifdial Organ Spec., Inc.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1980-1980

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Device record: <https://www.510kdatabase.net/k801632/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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