

K802956 GAMBRO A-V FISTULA NEEDLE SETDec 22, 1980
32 days to decisionK802956 · Product code: **FIE** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k802956/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	Nov 20, 1980
Decision date	Dec 22, 1980
Days to decision	32 days
Third-party review	No

APPLICANT

Company	Gambro, Inc.
Location	Mchenry, IL, US
510(k) history	86 submissions · 86 cleared · 1976-2009

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Device record: <https://www.510kdatabase.net/k802956/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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