

K013384 SHARGHP A.V. FISTULA NEEDLE SETJan 7, 2002
87 days to decisionK013384 · Product code: **FIE** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k013384/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	Oct 12, 2001
Decision date	Jan 7, 2002
Days to decision	87 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Shargh Pharmaceutical Co., Inc.
Location	Culver City, CA, US
Contact	ARMAND HAMEDANI
510(k) history	5 submissions · 5 cleared · 1990-2002

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