

K761098 FILTER DEVICESJan 28, 1977
67 days to decisionK761098 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k761098/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Nov 22, 1976
Decision date	Jan 28, 1977
Days to decision	67 days
Third-party review	No

APPLICANT

Company	Arrow Intl., Inc.
Location	Mchenry, IL, US
510(k) history	110 submissions · 105 cleared · 1976-2010

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Device record: <https://www.510kdatabase.net/k761098/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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