FDA 510(k) Premarket Notification

K761098 FILTER DEVICES

CLEARED

Jan 28, 1977
67 days to decision

K761098 · Product code: **FMI** · General Hospital
Source: 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