FDA 510(k) Premarket Notification

K781978 PUNCTURE KIT, DISPOSALBE VEIN

CLEARED

Dec 12, 1978
15 days to decision

K781978 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k781978/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Nov 27, 1978
Decision date	Dec 12, 1978
Days to decision	15 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k781978/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 22, 2026