FDA 510(k) Premarket Notification

K880350 YUE/FA DISPOSABLE SYRINGE

CLEARED

Feb 16, 1988
21 days to decision

K880350 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k880350/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jan 26, 1988
Decision date	Feb 16, 1988
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**Honway International Co.**
Location	San Francisco, CA, US
Contact	KEN CHIN
510(k) history	1 submissions · 1 cleared · 1988-1988