

**K880350 YUE/FA DISPOSABLE SYRINGE**Feb 16, 1988  
21 days to decisionK880350 · Product code: **FMF** · General HospitalSource: <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	<b>Honway International Co.</b>
Location	San Francisco, CA, US
Contact	KEN CHIN
510(k) history	1 submissions · 1 cleared · 1988-1988

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880350/>, Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 9, 2026