

**K000572 E.N.S.I. SYRINGE**Feb 24, 2000  
2 days to decisionK000572 · Product code: **MEG** · General Hospital  
Source: <https://www.510kdatabase.net/k000572/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Antistick (MEG)
Date received	Feb 22, 2000
Decision date	Feb 24, 2000
Days to decision	2 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Tri-Ject Intl. Corp.</b>
Location	Puyallup, WA, US
Contact	PETER G LEMIN
510(k) history	1 submissions · 1 cleared · 2000-2000

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k000572/> 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 4, 2026