FDA 510(k) Premarket Notification

K000572 E.N.S.I. SYRINGE

CLEARED

Feb 24, 2000
2 days to decision

K000572 · 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	**Tri-Ject Intl. Corp.**
Location	Puyallup, WA, US
Contact	PETER G LEMIN
510(k) history	1 submissions · 1 cleared · 2000-2000