

**K011283 MONOJECT PREFILL 0.9% SODIUM CHLORIDE FLUSH  
SYRINGE**Oct 15, 2001  
171 days to decisionK011283 · Product code: **NGT** · General Hospital  
Source: <https://www.510kdatabase.net/k011283/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Saline, Vascular Access Flush (NGT)
Date received	Apr 27, 2001
Decision date	Oct 15, 2001
Days to decision	171 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>The Kendall Company</b>
Location	Mansfield, MA, US
Contact	DAVID A OLSON
510(k) history	8 submissions · 8 cleared · 2001-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k011283/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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