

**K022090 SYNTEX PRE-POWDERED NITRILE EXAMINATION GLOVE**

Sep 3, 2002  
68 days to decision

K022090 · Product code: **LZA** · General Hospital  
Source: <https://www.510kdatabase.net/k022090/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Polymer Patient Examination Glove (LZA)
Date received	Jun 27, 2002
Decision date	Sep 3, 2002
Days to decision	68 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Syntex Healthcare Products Co., Ltd.</b>
Location	Xinji City, Hebei Province, CN
Contact	TAN SWU CHOON
510(k) history	17 submissions · 17 cleared · 2002-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k022090/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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