FDA 510(k) Premarket Notification

K022090 SYNTEX PRE-POWDERED NITRILE EXAMINATION GLOVE

CLEARED

Sep 3, 2002
68 days to decision

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

SUBMISSION DETAILS

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

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