FDA 510(k) Premarket Notification

K895023 LATEX PATIENT EXAMINATION GLOVE

CLEARED

Nov 16, 1989
119 days to decision

K895023 · Product code: **LYY** · General Hospital
Source: https://www.510kdatabase.net/k895023/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Jul 20, 1989
Decision date	Nov 16, 1989
Days to decision	119 days
Third-party review	No

APPLICANT

Company	**Beijing Latex Factory**
Location	Wilmington, MA, US
Contact	WEI T SHA
510(k) history	1 submissions · 1 cleared · 1989-1989