FDA 510(k) Premarket Notification

K891423 PATIENT EXAMINATION GLOVES (LATEX)

CLEARED

May 12, 1989
59 days to decision

K891423 · Product code: **LYY** · General Hospital
Source: https://www.510kdatabase.net/k891423/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Mar 14, 1989
Decision date	May 12, 1989
Days to decision	59 days
Third-party review	No

APPLICANT

Company	**J'Joy Intl., Inc.**
Location	Santa Fe Springs, CA, US
Contact	ROBERT WANG
510(k) history	1 submissions · 1 cleared · 1989-1989

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k891423/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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