FDA 510(k) Premarket Notification

K892131 PATIENT EXAMINATION GLOVES (LATEX)

CLEARED

Jul 14, 1989
101 days to decision

K892131 · Product code: **LYY** · General Hospital
Source: https://www.510kdatabase.net/k892131/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Apr 4, 1989
Decision date	Jul 14, 1989
Days to decision	101 days
Third-party review	No

APPLICANT

Company	**Five Lakes Co., Inc.**
Location	Taipei, TW
Contact	PHILIP LIN
510(k) history	1 submissions · 1 cleared · 1989-1989

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k892131/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 8, 2026