FDA 510(k) Premarket Notification

K892302 NON-STERILE PATIENT EXAMINATION GLOVES

CLEARED

Apr 28, 1989
22 days to decision

K892302 · Product code: **LYY** · General Hospital
Source: https://www.510kdatabase.net/k892302/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Apr 6, 1989
Decision date	Apr 28, 1989
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Yeou-Fwu Rubber Industrial Co., Ltd.**
Location	Shiang,Chang Huahsie, TW
Contact	WAN-FWU CHANG
510(k) history	2 submissions · 2 cleared · 1989-1989

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k892302/ 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