FDA 510(k) Premarket Notification

K892761 PATIENT EXAMINATION GLOVES (LATEX)

CLEARED

May 25, 1989
44 days to decision

K892761 · Product code: **LYY** · General Hospital
Source: https://www.510kdatabase.net/k892761/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Apr 11, 1989
Decision date	May 25, 1989
Days to decision	44 days
Third-party review	No

APPLICANT

Company	**E-Full Industrial , Ltd.**
Location	Kowloon Bay, HK
Contact	ELAINE CHIANG
510(k) history	2 submissions · 2 cleared · 1989-1989

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k892761/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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