FDA 510(k) Premarket Notification

K891881 PATIENT EXAMINATION GLOVE (LATEX)

CLEARED

May 25, 1989
58 days to decision

K891881 · Product code: **LYY** · General Hospital
Source: https://www.510kdatabase.net/k891881/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Mar 28, 1989
Decision date	May 25, 1989
Days to decision	58 days
Third-party review	No

APPLICANT

Company	**Cymun Development Sdn Bhd**
Location	Perak, Malaysia, MY
Contact	HOONG MUN
510(k) history	1 submissions · 1 cleared · 1989-1989

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k891881/ 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