FDA 510(k) Premarket Notification

CLEARED

K892267 PATIENT EXAMINATION GLOVES (LATEX)

Jul 14, 1989
100 days to decision

K892267 · Product code: **LYY** · General Hospital
Source: https://www.510kdatabase.net/k892267/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Apr 5, 1989
Decision date	Jul 14, 1989
Days to decision	100 days
Third-party review	No

APPLICANT

Company	**Ejs Sdn. Bhd.**
Location	Malaysia, MY
Contact	TAT ENG
510(k) history	1 submissions · 1 cleared · 1989-1989

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k892267/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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