FDA 510(k) Premarket Notification

K891661 PATIENT EXAMINATION GLOVES (LATEX)

CLEARED

May 25, 1989
66 days to decision

K891661 · Product code: **LYY** · General Hospital
Source: https://www.510kdatabase.net/k891661/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Mar 20, 1989
Decision date	May 25, 1989
Days to decision	66 days
Third-party review	No

APPLICANT

Company	**Dmp, Inc.**
Location	Houston, TX, US
Contact	JOY CHEW
510(k) history	3 submissions · 3 cleared · 1989-1989

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k891661/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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