FDA 510(k) Premarket Notification

K891863 PATIENT EXAM GLOVE (MFG. DALIAN LATEX & JIN ZHOU)

CLEARED

May 31, 1989
65 days to decision

K891863 · Product code: **LYY** · General Hospital
Source: https://www.510kdatabase.net/k891863/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Mar 27, 1989
Decision date	May 31, 1989
Days to decision	65 days
Third-party review	No

APPLICANT

Company	**China Xinxing Corp. , Ltd.**
Location	Beijing, CN
Contact	JING XIA
510(k) history	3 submissions · 3 cleared · 1989-1989

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k891863/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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