FDA 510(k) Premarket Notification

K902181 MODIFIED NON-STERILE LATEX EXAMINATION GLOVES

CLEARED

Jun 1, 1990
30 days to decision

K902181 · Product code: **LYY** · General Hospital
Source: https://www.510kdatabase.net/k902181/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	May 2, 1990
Decision date	Jun 1, 1990
Days to decision	30 days
Third-party review	No

APPLICANT

Company	**Aladan Corp.**
Location	Jessup, MD, US
Contact	BRADLEY L PUGH
510(k) history	18 submissions · 18 cleared · 1989-1996