

**K902181 MODIFIED NON-STERILE LATEX EXAMINATION GLOVES**Jun 1, 1990  
30 days to decisionK902181 · 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	<b>Aladan Corp.</b>
Location	Jessup, MD, US
Contact	BRADLEY L PUGH
510(k) history	18 submissions · 18 cleared · 1989-1996

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902181/> 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 June 29, 2026