

K913766 ANSELL NO POWDER AMBI PATIENT EXAMINATION GLOVESMar 11, 1993
567 days to decisionK913766 · Product code: LYY · General Hospital
Source: <https://www.510kdatabase.net/k913766/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Aug 22, 1991
Decision date	Mar 11, 1993
Days to decision	567 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Ansell, Inc.
Location	Dothan, AL, US
Contact	JOHN W MOUSHALL
510(k) history	30 submissions · 30 cleared · 1980-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k913766/> 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