FDA 510(k) Premarket Notification

K913766 ANSELL NO POWDER AMBI PATIENT EXAMINATION GLOVES

CLEARED

Mar 11, 1993
567 days to decision

K913766 · 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