FDA 510(k) Premarket Notification

K902282 ULTRA.SPEC NON-STERILE LATEX EXAMINATION GLOVE

CLEARED

Jun 7, 1990
16 days to decision

K902282 · Product code: **LYY** · General Hospital
Source: https://www.510kdatabase.net/k902282/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	May 22, 1990
Decision date	Jun 7, 1990
Days to decision	16 days
Third-party review	No

APPLICANT

Company	**Ncl Corp.**
Location	Akron, OH, US
Contact	BUSTER, SR.
510(k) history	1 submissions · 1 cleared · 1990-1990