

**K902282 ULTRA.SPEC NON-STERILE LATEX EXAMINATION
GLOVE**Jun 7, 1990
16 days to decisionK902282 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k902282/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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