

K013793 POWDER FREE LATEX EXAMINATION GLOVES WITH ALOE VERA @PH 5.5 AND PROTEIN LABELING CLAIMS

Jan 22, 2002
69 days to decision

K013793 · Product code: LYY · General Hospital
Source: <https://www.510kdatabase.net/k013793/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Nov 14, 2001
Decision date	Jan 22, 2002
Days to decision	69 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Shen Wei (Usa), Inc.
Location	Foster City, CA, US
Contact	BELLE L CHOU
510(k) history	58 submissions · 58 cleared · 1989-2022

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k013793/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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