FDA 510(k) Premarket Notification

K892579 ANATOMICAL EXAMINATION GLOVES

CLEARED

Jul 14, 1989
95 days to decision

K892579 · Product code: **LYY** · General Hospital
Source: https://www.510kdatabase.net/k892579/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Apr 10, 1989
Decision date	Jul 14, 1989
Days to decision	95 days
Third-party review	No

APPLICANT

Company	**Ansell Pacific, Inc.**
Location	Tucson, AZ, US
Contact	FRED DIETSCH
510(k) history	2 submissions · 2 cleared · 1989-1989

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k892579/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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