

**K892033 PATIENT EXAMINATION GLOVE (LATEX)**Aug 10, 1989  
132 days to decisionK892033 · Product code: LYY · General Hospital  
Source: <https://www.510kdatabase.net/k892033/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Mar 31, 1989
Decision date	Aug 10, 1989
Days to decision	132 days
Third-party review	No

**APPLICANT**

---

Company	<b>Weiss Ent., Inc.</b>
Location	Santa Rosa, CA, US
Contact	ROBERT WEISS
510(k) history	1 submissions · 1 cleared · 1989-1989

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k892033/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 8, 2026