

**K080606 ALATECH POWDERED LATEX PATIENT
EXAMINATION GLOVE**Aug 7, 2008
156 days to decisionK080606 · Product code: LYY · General Hospital
Source: <https://www.510kdatabase.net/k080606/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Latex Patient Examination Glove (LYY)
Date received	Mar 4, 2008
Decision date	Aug 7, 2008
Days to decision	156 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Alatech Healthcare, LLC
Location	Eufaula, AL, US
Contact	NEIL ANDERSON
510(k) history	4 submissions · 4 cleared · 1999-2009

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k080606/> Data sourced from FDA 510(k) public records (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. - Generated June 22, 2026