FDA 510(k) Premarket Notification

K080606 ALATECH POWDERED LATEX PATIENT EXAMINATION GLOVE

CLEARED

Aug 7, 2008
156 days to decision

K080606 · 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