FDA 510(k) Premarket Notification

K070760 POWDER FREE CHLORINATED LATEX PATIENT EXAMINATION GLOVE

CLEARED

Jun 7, 2007
79 days to decision

K070760 · Product code: **LYY** · General Hospital
Source: https://www.510kdatabase.net/k070760/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Mar 20, 2007
Decision date	Jun 7, 2007
Days to decision	79 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Derma Care Plus Products, LLC**
Location	Green Bay, WI, US
Contact	JOSEPH NEUSER
510(k) history	2 submissions · 2 cleared · 2007-2011