

**K070760 POWDER FREE CHLORINATED LATEX PATIENT EXAMINATION GLOVE**Jun 7, 2007  
79 days to decisionK070760 · 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        | <b>Derma Care Plus Products, LLC</b>  |
| Location       | Green Bay, WI, US                     |
| Contact        | JOSEPH NEUSER                         |
| 510(k) history | 2 submissions · 2 cleared · 2007-2011 |

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k070760/> 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 18, 2026