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Omni's Family of Gloves

- 111 Series Latex Powdered Examination Gloves
- 112 Series Latex Powder Free Examination Gloves
- 113 Series Latex Powder Free Examination Gloves
- 202 Series "Lite Touch" Nitrile Examination Gloves
- 212 Series Nitrile Examination Gloves For EMS & Chemotherapy
- 311 Series Powdered Vinyl Examination Gloves
- 312 Series Powder Free Vinyl Examination Gloves
- 361 Series Powdered NSF Food Service Vinyl Gloves
- 362 Series Powder Free NSF Food Service vinyl Gloves
- 412 Series Powder Free Stretch Vinyl Examination Gloves
- 131 Series Powdered Latex Surgeons Gloves
- 132 Series Powder Free Latex Surgeons Gloves

GLOVE LINES

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e-mail addresses to
News@OmniGloves.com
and we will send them
our next issue.



OMNI Leads in Vinyl Glove Production!

Omni International is well connected in China. Our primary vinyl producer is a 100% owned subsidiary of a major Chemical Corporation, the second largest chemical company in China. They, in fact, provide chemicals to the majority of glove manufacturers located throughout China.

Omni began our business relationship with our manufacturing partner in 2003. We were their first U.S. account and actually helped them with all of their FDA filings as well as assisted them in setting up their QA and QC procedures.

Today, they are one of the largest vinyl glove producers in China. We are in an enviable position of not only having one of the premier glove manufacturers as our supplier, but we also have a very strong friendship which has been built up by really being partners.

In December of 2008, the new regulations will be put into effect by the FDA changing acceptable AQL levels for ALL Medical Gloves from 4.0 to 2.5. These numbers are sometimes confusing to the consumer. All factories use statistical sampling to assure that gloves shipped to the U.S. meet the current AQL requirements. As you may be aware of, the FDA randomly pulls containers, samples their content and sends these samples to an FDA laboratory

for analysis. Should these gloves fail in any way due to pin holes, the entire lot (in most cases the entire container) must be either destroyed or returned to China.

In an effort to maintain the highest quality standard and to prevent having any FDA failures, Omni, as well as other high quality glove manufacturers, release product based on a more stringent sampling plan. Omni and our partner state all of our gloves must pass a 1.5 AQL inspection before they are released for shipment to the U.S. Utilizing this higher AQL level produces a higher quality of product, free of manufacturing defects and also assures the highest quality standard that can easily pass any FDA testing.

As we stated in our bulletin, **Market Outlook for 2008** (available from Omni), there are going to be many glove manufacturers that will not be able to meet this new criteria. This is due to several issues. First and foremost, many producers do not have state of the art manufacturing facilities. Secondly, they do not have the necessary technical personnel and thirdly, they do not have the processes in place to produce this level of higher quality products.

Rest assured, Omni and our manufacturing partners are absolutely ready to surpass all expectations. For any and all questions contact us at 888-999-6664.

NEED SAMPLES? Please contact us at News@OmniGloves.com or call 888-999-6664. **THANK YOU** for buying Omni Glove products.