



December 1, 2012

To Whom It May Concern,

This will confirm that the US Food and Drug Administration regularly inspects Kutol Products Company where we manufacture "Human Topical" Over The Counter (OTC) Drug Products. While the FDA does not "approve" manufacturers or products they do make sure that producers and packers comply with the applicable regulations.

Kutol's manufacturing facility was last inspected by the FDA in the Fall of 2012. The FDA inspects Kutol according to GMP standards as well as compliance to specific regulations for OTC products. Based on our last inspection we remain in good standing with the FDA concerning our level of compliance with both GMP and more specific OTC regulations.

**Our manufacturing site registration number is 1513988 and our FDA labeler code # is 50865.**

A small sampling of OTC Items manufactured by Kutol include:

5666 Instant Hand Sanitizer 12/800ML

5619 Instant Hand Sanitizer 12/8OZ

5019 Antibac Hand Soap 12/8OZ

68841 Alcohol Hand Sanitizer 6/1000ML

64041 Antibacterial Moisture Wash 6/1000ML

A7802F TidyFoam Antibacterial Soap 4/1000ML

Please contact me if you have any questions on this matter or desire to know more about how we can take care of your and hygiene needs.

Sincerely,

KUTOL PRODUCTS

A handwritten signature in black ink that reads "Bob Bennett". The signature is written in a cursive, flowing style.

Vice President, Sales