

What process does N2G use to determine product safety and the highest level of quality?

N2G uses a 3-Phase Testing Protocol to ensure the safest and highest quality products are delivered to you

Phase 1: Pre-Production - Incoming Raw Material Testing

Identity: To confirm the ID of a given raw material and the accuracy of the Certificate of Analysis (COA) we use FT-NIR (fourier transform near-infrared technology).

Purity: Vendor COA claims are checked for purity by HPLC, GC or ICP.

Heavy Metals: We test for arsenic, cadmium, lead and mercury. Heavy metals must be under the USP specified guidelines.

Microbial: We test for total plate count, total yeast and mold count and Coliform bacteria to make sure that the raw material complies with the USP, AHPA and FDA guidelines.

Pathogens: Raw materials are tested for E. coli, Salmonella, and Staphylococcus aureus pathogens (as needed).

Drug Screen: Raw materials are tested for banned steroidal substances (as needed)



Phase 2: Intra-Production Testing

Cleanliness: We perform ATP swab tests prior to batch production and allergen swab tests as needed to ensure equipment is free of potential contamination or microbes.

Identity Check: We use FT-NIR to confirm the ID of a given in-process mix to ensure it contains the correct raw ingredients.

Purity Check: Purity is checked by using HPLC, GC, and/or ICP to ensure the in-process mix will create a final product that meets label claims.

Phase 3: Post-Production Testing

Identity Check: We use FT-NIR to confirm the ID of a finished product.

Purity (LABEL CLAIM): We check the purity by HPLC, GC, and/or ICP if there is a label claim on the finished product COA.

Microbial Check: We test for total plate count, total yeast and mold count and Coliform bacteria to make sure that the finished product complies with specifications.

Pathogens Check: We test again for the following pathogens, E. coli, Salmonella, and Staphylococcus aureus (as needed).

What Does This All Mean?

100% of all inbound raw material will be FDA validated

100% of all finished products will be FDA validated

100% of finished products will always meet label claims

Now that you truly understand what it takes to offer this level of quality, can you really afford to put your clients on any other products?