Retained samples of all raw materials and finished goods are maintained for one year past expiration date.

Accurate formulation of the nutritional ingredient systems should take into account potency losses which can occur due to processing, packaging, and shelf life stability issues.

Incoming raw materials should meet FCC or USP/NF where such specifications are listed and should be quarantined upon receipt.

Materials released for use in production by QC approval.

Validated analytical test methods are employed for creation of a Certificate of Analysis on finished goods.

Commitment to cGMP standards & commitment to quality in all processes.

Science and Technology: Which additional technologies are available in-house:

- Wet/Dry Blending
- Microencapsulation
- Trituration
- Spray Drying
- Hot Melt Granulation
- Agglomeration
- Grinding & Micronizing
- Instantizing