

June 25, 2008 FDA Deadline For Dietary Supplement Compliance With Good Manufacturing Practice (GMP) Regulation

Dietary Supplement manufacturers with 500 or more employees are required to be in compliance with GMP regulation by June 25, 2008. Manufacturers with less than 500 employees, but more than 20 have to comply by June 2009. It is anticipated that many retailers will expect their suppliers to demonstrate evidence of diligent progress toward compliance in advance of these regulatory deadlines. [FDA Final Rule cGMP Requirement](#).

Shuster Laboratories, an STR Company, offers comprehensive GMP regulatory compliance services including:

- Facility audits to verify GMP compliance.
- Analytical and microbiological testing on raw materials and finished products to ensure GMP compliance for identity, strength, composition, potency, purity and stability. These services are available at our Canton, MA and Shanghai, China laboratories.
- Contaminant testing capabilities.
- Method development/qualification for raw materials and finished product testing.
- Consulting services on rule interpretation, label review and gap analysis specific to the new GMP regulations.

We can help ensure your company has the highest level of confidence in its products, raw materials, and manufacturing practices. For more information, please contact Dr. Robert Rolle at 800-444-8705, ext. 140 or at f.robert.rolle@shusterlabs.com.