The dietary supplement industry is becoming increasingly regulated, and if you're not careful, actions taken by your CMO can get your product recalled by the FDA. Knowing your responsibilities, what quality control tests you or your CMO should be performing, and what analytical techniques to use will make sure you’re in full compliance with today’s cGMPs.

**INDUSTRY GROWTH MEANS REGULATION**

But new regulations bring new complexities.

**The dietary supplements industry is now the largest industry in the state of Utah.**

**$7 BILLION IN UTAH ALONE**

**HIGHLY REGULATED INDUSTRY**

**CONSUMER AND PRACTITIONER CONFIDENCE**

**MORE SALES**

**GROWTH**

**FDA OVERSIGHT**

If you create dietary supplements, using a CMO to manufacture your products does not relieve you of any compliance requirements under 21CFR part 111.

**CMO**

**PACKER**

**CONTRACT TESTING LAB**

**FULFILLMENT SERVICE**

**REGULATORY AFFAIRS CONSULTANT**

**SHIPPER**

**DISTRIBUTOR**

**YOU ARE LIABLE FOR THE CGMPs OF EVERY**

**SEPARATION SCIENCE**

Chromatography plays a vital role in meeting the numerous regulatory compliance requirements laid out in the cGMPs, and no one technique or system can perform all of the required testing needed for compliance.

**COMPLIANCE AREAS**

**IDENTITY**

100% of all supplements must be identity tested

**RACs**

Ensure freedom from reasonably anticipated contaminants

**RAW MATERIALS**

Ensure potency of raw material claims

**FINISHED PRODUCTS**

Ensure label claim is satisfied

**Quality Control of Dietary Supplements: Chromatographic Approaches for cGMP Compliance**

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**INTERESTED IN LEARNING MORE?**

Sign up for the webcast at www.chromatographyonline.com/quality_control