FOOD/BEVERAGE & CHEMICAL TOXICOLOGY

Regulatory Services and Ingredient Approval

Trinity Consultants is a leading global environmental consulting firm that brings 50 years of experience providing services and solutions in the EHS Regulatory Compliance, Built Environment, Life Sciences, and Water & Ecology markets. Trinity has the technical expertise and acute industry knowledge to provide optimal solutions to highly complex pharmaceutical development, process, safety, and facility design challenges.

SafeBridge, a Trinity Consultants team, has an experienced staff of toxicologists and registered quality assurance professionals (RQAP-GLP) to assist clients with regulatory approvals for biocides, specialty chemicals, food and beverage ingredients, food contact substances, and dietary supplements. Our extensive experience and highly organized program management enables us to shorten the timeframe typically associated with regulatory approvals, often compressing time to market. Of critical importance, we have a track record of constructive interface with regulatory agency contacts in the U.S., Canada, Europe, and other countries on scientific and technical issues that benefit our clients. We are especially effective at building novel and successful strategies to ensure that our clients are in the strongest regulatory and safety positions possible. In many cases, our guidance and/or interaction with regulatory agencies has resulted in the most effective use of resources for ingredient approval or product registration.

Our services include the following:

- **Regulatory Strategy Development:** Identifying, interpreting, and applying regulations and policy
- **Literature Searching and Reviews:** Systematic and targeted literature searches and reviews to support regulatory activities
- Data Interpretation: Evaluating data and information to perform data gap analysis and safety assessments, read-across, and/or QSAR assessments
- **Safety Testing:** Design, implementation, and management of toxicology testing programs conducted according to Good Laboratory Practice (GLP) standards
- **Quality Assurance (QA) and GLP Support:** Quality assurance oversight of GLP testing programs including test guideline compliance, study audits, and scientific/technical reviews of study reports

- **Dossier Development:** Preparation of application dossiers for food additives, flavoring ingredients, Generally Recognized as Safe (GRAS) materials, food contact substances, and New Dietary Ingredients, and preparation of waiver and bridging justifications for FIFRA data requirements
- **Develop Exposure and Intake Assessments:** Exposure/intake assessments and calculations supporting risk assessments and regulatory dossiers
- **Coordinate Expert Panels:** Recruit, manage, and facilitate Expert Panels addressing scientific questions and regulatory requests
- **Scientific and Regulatory Affairs:** Provide understanding of safety and regulatory risks associated with finished products, ingredients, or product constituents of interest, and direct engagement with regulatory authorities

CONTACT OUR TEAM!

For more information about how we can help your organization, please contact us.

Trinity Consultants | SafeBridge
P 650.961.4820 / *info@safebridge.com*

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