



REACH Consulting Services

Regulatory and Scientific Specialists

Support for Pre-Registration, Registration and Risk Assessment

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REACH 2018 registration deadline has passed, but are you compliant? CSI can help!

The REACH Regulation has presented a unique challenge to the chemical industry throughout the world. CSI has been actively involved with REACH since before the Regulation came into force in 2007, and offers a comprehensive service to companies and consortia facing up to its demands.

CSI's team of regulatory scientists has accumulated a wealth of experience in dealing with all scientific and regulatory aspects of REACH, and specialises in the provision of technical and administrative support for compliance with the REACH Regulation. We are therefore ideally equipped to assist our clients in ensuring that they meet their REACH obligations in a timely and cost effective manner. In addition to direct scientific and regulatory support, CSI offers Only Representative (for non-EU companies) and Third Party Representative services.

Compliance Services International (CSI) provides comprehensive support and guidance to assist with all aspects of REACH compliance, from general consultancy advice to complete dossier compilation and submission.

Only Representative (OR) / Third Party Representative

Only Representative role for non-EU companies, independent Third Party support for handling confidential information. QSAR/structural read-across proposals, and submission of pre-registration information.

Analytical Data Support

Providing adequate evidence of substance identity is a fundamental part of the registration dossier for joint submission members. CSI can advise on specific substance identity requirements, review existing analytical data for completeness, and place and monitor analytical programmes at specialist testing facilities.

Chemical Safety Assessments (CSA) / Chemical Safety Reports (CSR)

Hazard assessment, exposure assessment, and risk characterization. Review of existing data, identification of data gaps, development of testing strategies. Placement and monitoring of studies at contract research organizations.

Registration Dossier Submission

Preparation of technical dossiers, safety data sheets (SDS), chemical safety assessments (CSA), chemical safety reports (CSR) and supporting IUCLID databases. Submission of documentation to the European Chemicals Agency (ECHA).