Intertek Health Environmental & Regulatory Services (HERS) supports its customers with the technical know-how needed to create robust dossiers and chemical safety reports, ensuring a company can meet this final 2018 REACH deadline, by doing the right thing, the right way, the first time.

Background
European Union (EU) manufacturers and importers need to prepare for the final REACH registration deadline on the 1st June 2018, which covers all phase-in substances manufactured or imported into the EU at and above 1 tonne per year not previously registered.

Those in scope need to implement their action plans now, to ensure they will remain in the market. There are consequences to not meeting the deadline. Non-compliance with the REACH regulation could subsequently lead to the removal of substances from the EU market. This could result in not only financial costs to business but damage to brand and reputation with customers.

Many of those involved with 2010 and 2013 registrations will understand the importance of being well prepared for the fast approaching 2018 deadline and the need to register now. A company is exposed to increased business risk by starting compliance efforts late, and time required should not be underestimated.

Boost your stretched resources and/or avoid potentially costly errors made by the inexperienced and benefit from our experts and their deep knowledge of the REACH regulation.

Intertek has an enviable record of successful 2010 and 2013 registrations and can support your 2018 registration needs delivering comprehensive registration services for REACH in a timely and cost effective manner.

Advantages to Acting Now
• Management of registrations in the required timeframe
• Testing and studies completed in time
• Avoidance of compromises to meet compliance
• Enough time for reworking the dossier following submissions with updates

REACH Registration Technical Dossier Services
• REACH Technical Dossier data gap analysis:
  • Perform literature searches to locate available data on substances of interest
  • Evaluate existing study reports and rate according to Klimisch criteria
  • Analyse available data against the REACH requirements to identify key studies and any gaps
  • Confirm key studies and prepare robust and endpoint summaries for IUCLID
  • Develop Derived-No-Effect-Levels (DNELS)
  • Fill any gaps using scientific strategies that may include the use of:
    • REACH testing (substance identification, sameness testing, physicochemical property determination, toxicological, ecotoxicological, etc.)
    • Exposure Scenarios
    • Waiver arguments
    • (Q)SARs and extrapolation
    • Read-across data
  • Creation of IUCLID 5 Dossier
  • Preparing relevant Chemical Safety Reports (CSR) that cover client’s use patterns
  • Submission of completed REACH Registration Dossier to ECHA
  • SIEF and Consortia Management when needed
  • Review and revise classification and labelling (EU CLP / UN GHS) and prepare extended Safety Data Sheets
  • Consulting support from Intertek REACH Specialists
REACH Services

- Conduct an assessment of a company's current REACH status: review all substances and determine obligations under REACH, identify company obligations, gaps in information, provide a compliance strategy
- Act as an Only Representative for non-EU companies wishing to register their substances for their supply chain in the EU – includes annual reporting of volumes, provide analysis reports each year
- Provide reporting services for OR clients’ Non-EU customers through OR Supply Chain services
- Provide reporting services for clients’ supplier covered substances and collection/retention of Declaration of REACH Compliance (DRC)
- Provide annual reporting/analysis for Non-OR clients to monitor volumes for potential imminent registrations
- Provide assistance in building joint submission dossiers: communicate with SIEF, obtain LOA, act as a third party representative and develop substance identification and registration strategies
- Conduct analytical testing necessary to complete registration of substances

REACH Testing Programs to Meet the 2018 Deadline

Based on our extensive knowledge of global chemical regulatory issues and considerable experience gained during the previous deadlines, we continue to offer REACH testing programs for the 2018 deadline through tailored programs that are designed to facilitate your successful registration.

These programs include substance identification testing, sameness and physico-chemical property testing, delivering highly relevant technical guidance to ease your registration process. With strengths in data assessment, we can identify any data gaps in the information to complete a dossier and then put in place efficient strategies to address these gaps whilst minimizing unnecessary testing costs. Services can be delivered to ISO 17025 or Good Laboratory Practice (GLP) standards as required.

The Intertek Advantage

Intertek Health, Environmental & Regulatory Services (HERS) has extensive expertise and experience having worked with a variety of clients across multiple industries. Let our experienced professionals take care of your REACH obligations, absorbing this peak demand for regulatory resource and expertise, allowing your people to focus on core business needs. We can support your brand protection and market share through regulatory compliance. Client’s benefiting from our services can enjoy lower costs through outsourcing and continuous, convenient and accessible support.

About Intertek

Intertek is a leading Total Quality Assurance provider to industries worldwide. Our network of more than 1,000 laboratories and offices and over 42,000 people in more than 100 countries, delivers innovative and bespoke Assurance, Testing, Inspection and Certification solutions for our customers’ operations and supply chains. Intertek Total Quality Assurance expertise, delivered consistently with precision, pace and passion, enabling our customers to power ahead safely.

Total Quality. Assured.