



REACH out for chemicals in production

Manufacture of medicinal products involves a range of different substances such as precursors, intermediates, excipients, and process chemicals. *EC Regulation 1907/2006* on Registration, Evaluation, Authorisation and Restriction of Chemical substances (REACH) requires registration and information on the chemical safety for these substances to the European Chemicals Agency (ECHA). The information requirements depend on the quantity of the substance that is manufactured or imported into the EU. SAXOCON provides you with everything you need to document REACH compliance in accordance with regulatory requirements including:

- Gathering and review of existing toxicological data
- Data gap filling using (Q)SAR and read-across methods
- Submission of data via IUCLID

Why choose us?

SAXOCON services for manufacture of pharmaceutical products gives you access to:

- Toxicologists with experience from the Danish Environmental Protection Agency with extensive knowledge of REACH regulation and EU Classification and Labelling of chemical substances
- Best in class experience in using computational toxicology including (Q)SAR modelling and read-across methods
- Extensive experience in IUCLID submission and other REACH-IT solutions

Delivery

SAXOCON compiles a written report comprising the toxicological information necessary for fulfilling REACH requirements.