### In face of REACH

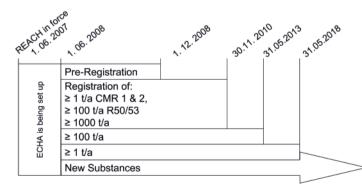
REACh ChemConsult GmbH and TCP Dr. Genz (HK) Co. Ltd.

help you to bring your products to the EU.

If you are a Non-EU manufacturer of substances on its own, in preparations and to a certain extent in articles, you should appoint an Only Representative in the EU in order to keep your existing distribution chain and to be flexible on your EU-costomers. After the preregistration of your substances the SIEF (substance information exchange forum) will form in order to share costs and data for the registration of the substances at the ECHA (European Chemicals Agency).

First necessary step:

Get an **Only Representative** in the EU for the pre-registration to benefit from the extented registration deadlines for the registration of the phase-in substances you market in the EU.



Pre-register phase-in substances to benefit from extended deadlines





### **REACH Services**

- Only Representative Services for
- Pre-Registration
- SIEF Participation
- Registration
- Seminars on roles and duties to make your company REACH compliant.

### **Contact**

# • in Europe (Germany):

Dr. Thomas Gildemeister REACh ChemConsult GmbH World Trade Center, Freiberger Str. 39,

D-01067 Dresden, Germany

Tel: +49 (0)351 476 930 0 Fax: +49 (0)351 476 930 15

Email: info@reach-chemconsult.com Website: www.reach-chemconsult.com

# • in ASIA (Hong Kong):

Mr. S. Postrach

TCP Dr. Genz (HK) Co. Ltd.

26/F., Tamson Plaza, 161 Wai Yip Street Kwun Tong,

Kowloon, Hong Kong

Tel: +852 2389 2200 Fax: +852 2389 3073

Email: tcp.hk@tcp-labs.com.hk

Website: www.tcp-labs.com

# **Our Expertise**

The experts of *REACh ChemConsult GmbH* combine a long time professional and scientific experience in chemical assessment, in developing, performing, and monitoring of (eco)-toxicity tests and in process risk assessment towards health, safety and environmental regulations and standards from activity at the chemical industries, university and governmental agencies.

#### **Chemical registration procedures**

- REACH
- Registration of new substances according to EC Directive 92/32/EEC
- Registration of human drugs according to EMEA Directives
- Chemical assessment within CICAD (WHO)
- Professional experience in the German national existing substance programme
- OECD ICCA HPV (High Production Volume programme) through activities at BUA<sup>1</sup>

#### Scientific research experience through

 participation in research projects of German EPA<sup>2</sup>, CEFIC<sup>3</sup> and industry

<sup>1</sup>BUA: Advisory Committee on Existing Chemicals of the German Chemical Society (GDCh), <sup>2</sup>EPA: environmental protection agency <sup>3</sup>CEFIC: European Chemical Industry Council

