REACH GUIDANCE DOCUMENT AVAILABILITY
As of July 30, 2007

REACH Guidance Document	Availability	Number of Pages	RIP Number[1]
Guidance mainly for Industry Use			
Guidance on registration Reference name: *Guidance on registration* This document describes when and how to register a substance under REACH. It consists of two parts: one on Registration tasks and obligations and the other on the preparation of the Registration Dossier.	Current	106	NA
Guidance on pre-registration Reference name: *Guidance on pre-registration* This document describes how to identify the substances that can be pre-registered as well as when and how to pre-register them.	Fall, 2007	NA	3.4
Guidance on data sharing Reference name: *Guidance on data sharing* This document describes data sharing mechanisms for phase-in and non phase-in substances under REACH. It includes the communication within the SIEF and the cost sharing guidance. The document also describes the Confidential Business Information and Competition Law issues in the context of data sharing.	Fall, 2007	NA	3.4
Guidance for intermediates Reference name: *Guidance for intermediates* This document describes when and how the specific provisions for the registration of intermediates under REACH can be used.	Current	27	NA
Guidance for monomers and polymers Reference name: *Guidance on polymers* This document describes the specific provisions for polymers and monomers under REACH.	Current	18	NA

[1] More information can be found on the status of REACH implementation projects (RIP) at http://ecb.jrc.it/reach/rip/

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Guidance on Scientific Research and Development (SR&D) and Product and Process Oriented Research and Development (PPORD) Reference name: *Guidance on PPORD* This document describes specific provisions under REACH for substances manufactured, imported or used in scientific Research and Development (SR&D) and Product and Process Oriented Research and Development (PPORD).	Current	17	NA
Guidance on Classification and Labelling notification Reference name: *Guidance on C&L notification* This document describes when and how to notify a classification and labeling for a substance under REACH.	2008	NA	Unknown
Guidance on requirements for substances in articles Reference name: *Guidance for articles* This document assists producers and importers of articles in identifying whether they have obligations under REACH; in particular in relation to registration and notification according to Article 7, and in relation to article supply chain communication according to Article 33.	Summer, 2007	NA	3.5
Guidance for Downstream Users Reference name: *Guidance for Downstream Users* This document describes the roles and obligations of downstream users, and advises them on how to prepare for the implementation for REACH.	Summer, 2007	NA	3.5

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Guidance on the preparation of an application for authorisation Reference name: *Guidance on authorisation application* This document describes how to prepare an application for authorisation and provides guidance on analysis of the alternatives and substitution plan. It also describes how third parties may prepare and submit information on alternatives.	Winter, 2007	NA	3.7
Guidance mainly for Authorities Use			
Guidance on Dossier and Substance Evaluation Reference name: *Guidance on evaluation* This document describes the evaluation tasks to be performed by the Authorities: evaluation of testing proposals and compliance check by the Agency and substance evaluation by the Member States Competent Authorities.	Current	139	NA
Guidance on Dossier and Substance Evaluation Reference name: *Guidance on evaluation* This document describes the evaluation tasks to be performed by the Authorities: evaluation of testing proposals and compliance check by the Agency and substance evaluation by the Member States Competent Authorities.	Current	60	NA
Guidance on Dossier and Substance Evaluation Reference name: *Guidance on evaluation* This document describes the evaluation tasks to be performed by the Authorities: evaluation of testing proposals and compliance check by the Agency and substance evaluation by the Member States Competent Authorities.	Current	58	NA

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REACH Guidance Document	Availability	Number of Pages	RIP Number[1]
Guidance on inclusion of substances in Annex XIV (substances subject to Authorisation) Reference name: *Guidance on Annex XIV inclusion* This document describes how the authorities (the Agency in co-operation with the Member States Competent Authorities) will include substances in the authorisation system. Guidance is given on the elaboration of the dossier that supplements each recommendation of a substance for inclusion in Annex XIV.	Fall, 2007	NA	4.3
Guidance for the preparation of an Annex XV dossier for restrictions Reference name: *Guidance on Annex XV for restrictions* This document describes how the authorities (Member States Competent Authorities or the Agency on request from the Commission) can prepare an Annex XV dossier to propose a restriction under REACH.	Current	130	NA
Guidance on the different methods under REACH			
Guidance for identification and naming of substances in REACH Reference name: *Guidance on substance identification* This document describes how to name and identify a substance under REACH.	Current	115	NA

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REACH Guidance Document	Availability	Number of Pages	RIP Number[1]
Guidance on how to comply with the provisions of the new Regulation on Classification, Packaging and Labelling of substances and mixtures Reference name: *Guidance on Classification, Packaging and Labelling* This document aims to assist industry and authorities to implement the new GHS criteria within the EU which are based on the UN Globally Harmonised System for the Classification and Labelling of chemicals (GHS) and to fulfil the relevant procedures.	2008	NA	3.6
Guidance for the preparation of the Chemical Safety Report Reference name: *Guidance on Chemical Safety Report* This document aims at assisting industry in conducting Chemical Safety Assessments and preparing Chemical Safety Reports, when required, as part of a registration dossier (for a substance on its own or as part of a preparation or as released from an article), as part of an authorisation application or as part of downstream user obligations. It also sets out the basic principles for authorities preparing a risk assessment in support of a restriction proposal, and when required as part of a Substance Evaluation.	2008	NA	3.2

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REACH Guidance Document	Availability	Number of Pages	RIP Number[1]
Guidance on information requirements under REACH Reference name: *Guidance on information requirements* This document provides guidance on the collection and assessment of the available information on the intrinsic properties of the substances to be registered, on the requirements specified by REACH, on the identification of data gaps and on the generation of the additional information required to comply with the Regulation.	Fall, 2007	NA	3.3
Guidance on Socio Economic Analysis Reference name: *Guidance on Socio Economic Analysis* This document assists the different actors in preparing a socio-economic analysis or input for one as part of the Authorisation and Restriction procedures.	2008	NA	3.9
Guidance on priority setting for evaluation Reference name: *Guidance on priority setting for evaluation* This document describes the different priority setting methods developed to prioritise dossiers, testing proposals or substances for evaluation and gives guidance for the Agency and the Member States Competent Authorities on the application of these methods.	Fall, 2007	NA	4.5
Guidance on IUCLID Reference name: *Guidance on IUCLID* This document describes how to use IUCLID 5 and how to prepare the dossiers for different REACH requirements.	Current	2044	NA