

ECHA Guidance for Downstream users The essentials

Communication in the supply chain
Chapters 1, 3 and 6

5 March 2014

Laura Walin
Directorate of Registration
European Chemicals Agency



Chapter 1: Introduction



- REACH processes
- Overview of communication in the supply chain
- Key terms

REACH – main processes from downstream user perspective



Registration

Communicate uses and conditions of use to registrants
Receive extended safety data sheets



Member States

Evaluation

- Dossier evaluation
- Substance evaluation

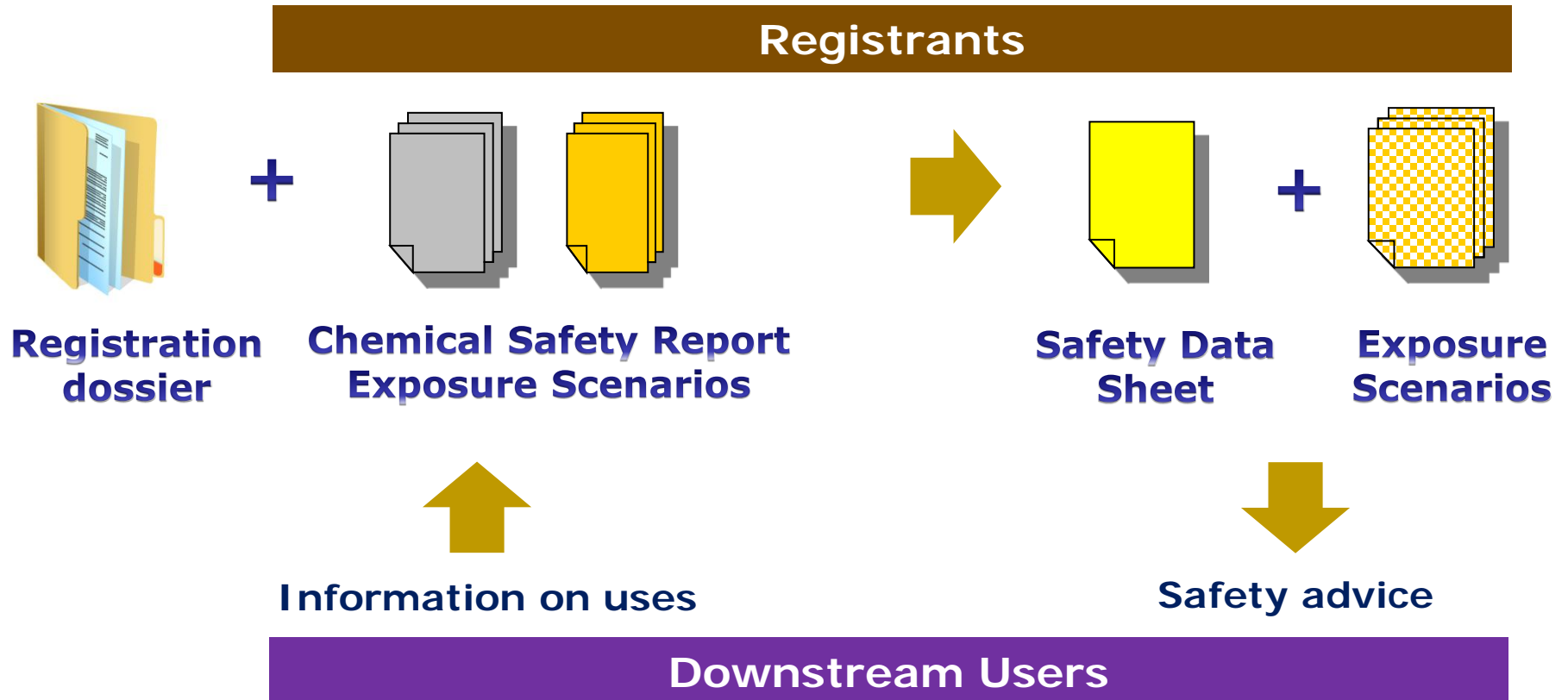
If registrant's chemical safety assessment changes, downstream user may receive an updated SDS
Substances may be identified as substances of very high concern

Authorisation Restriction

Downstream user's use needs to be covered by authorisation (own or upstream)
Only unrestricted uses may continue



Communication in the supply chain

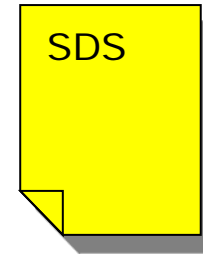


Key terminology

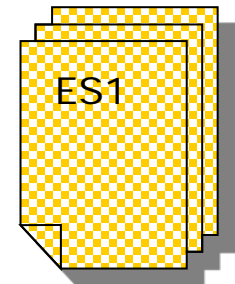
Use



Exposure scenario



+



Conditions of use

Operational conditions



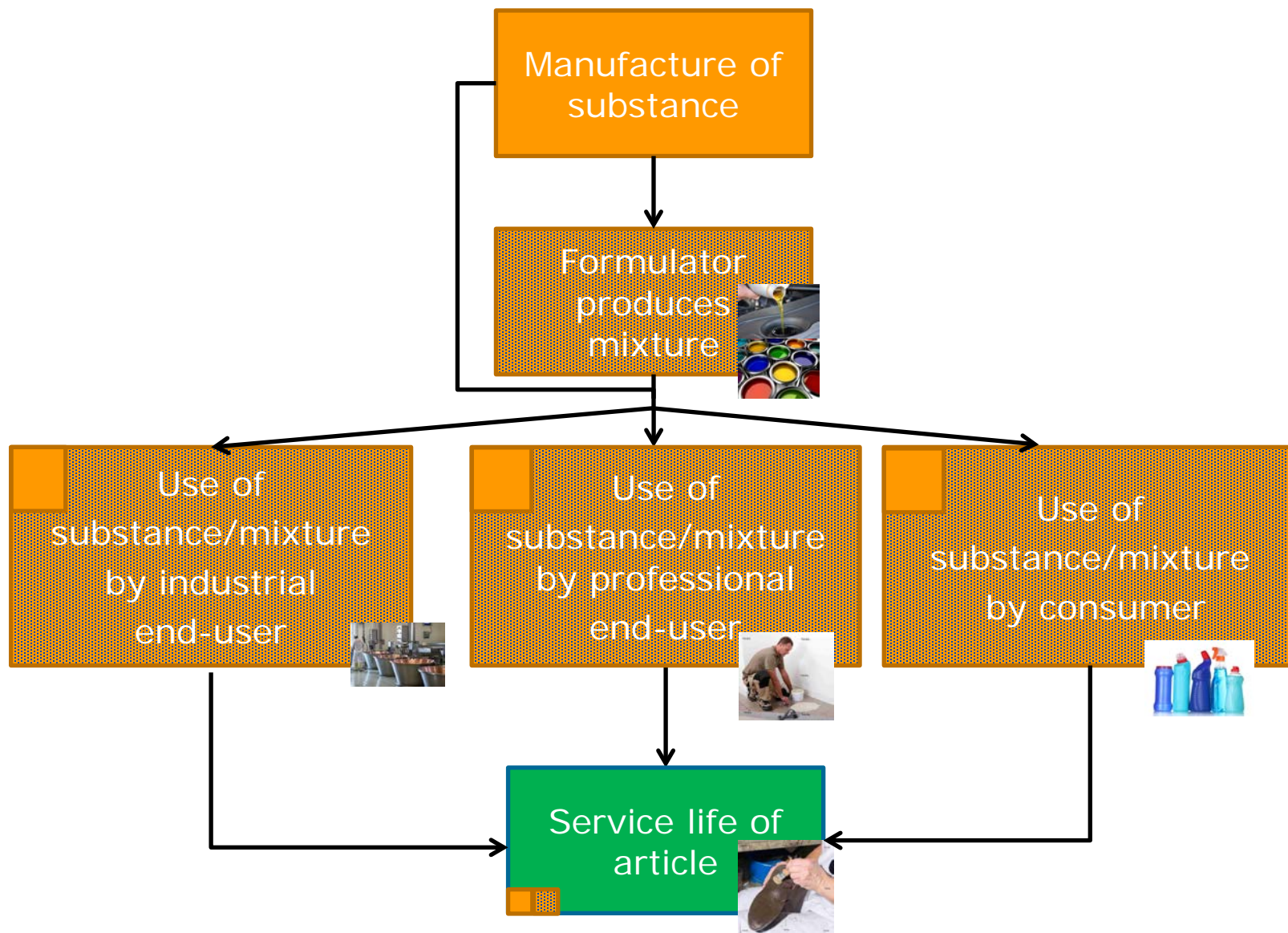
Risk management measures



Chapter 3: Communication in the supply chain

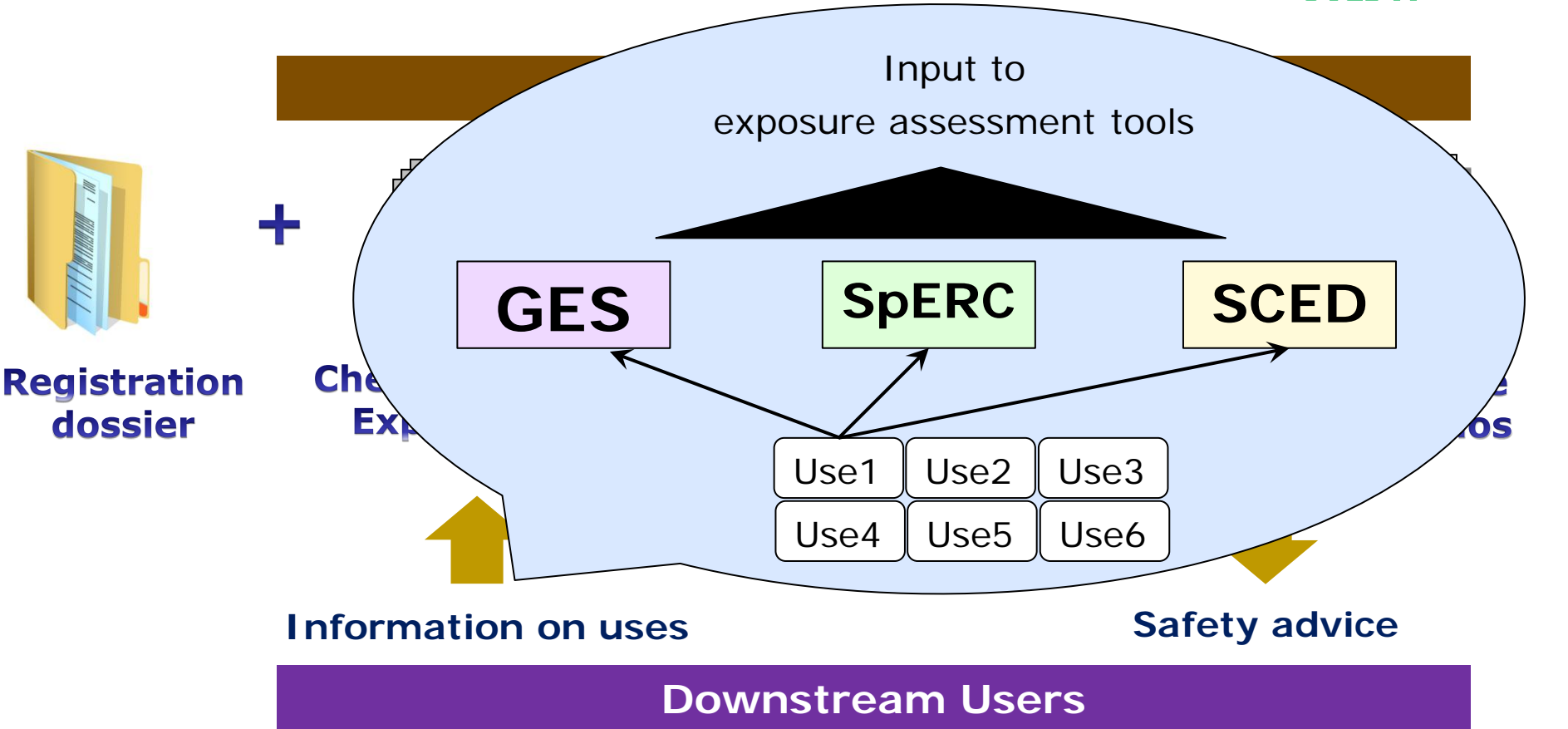


- Life cycle of a substance
- Harmonised communication elements **NEW**
- Communication via sector organisations **NEW**
- Communication from downstream user to supplier
- Response to downstream user communication



Harmonised communication elements

NEW

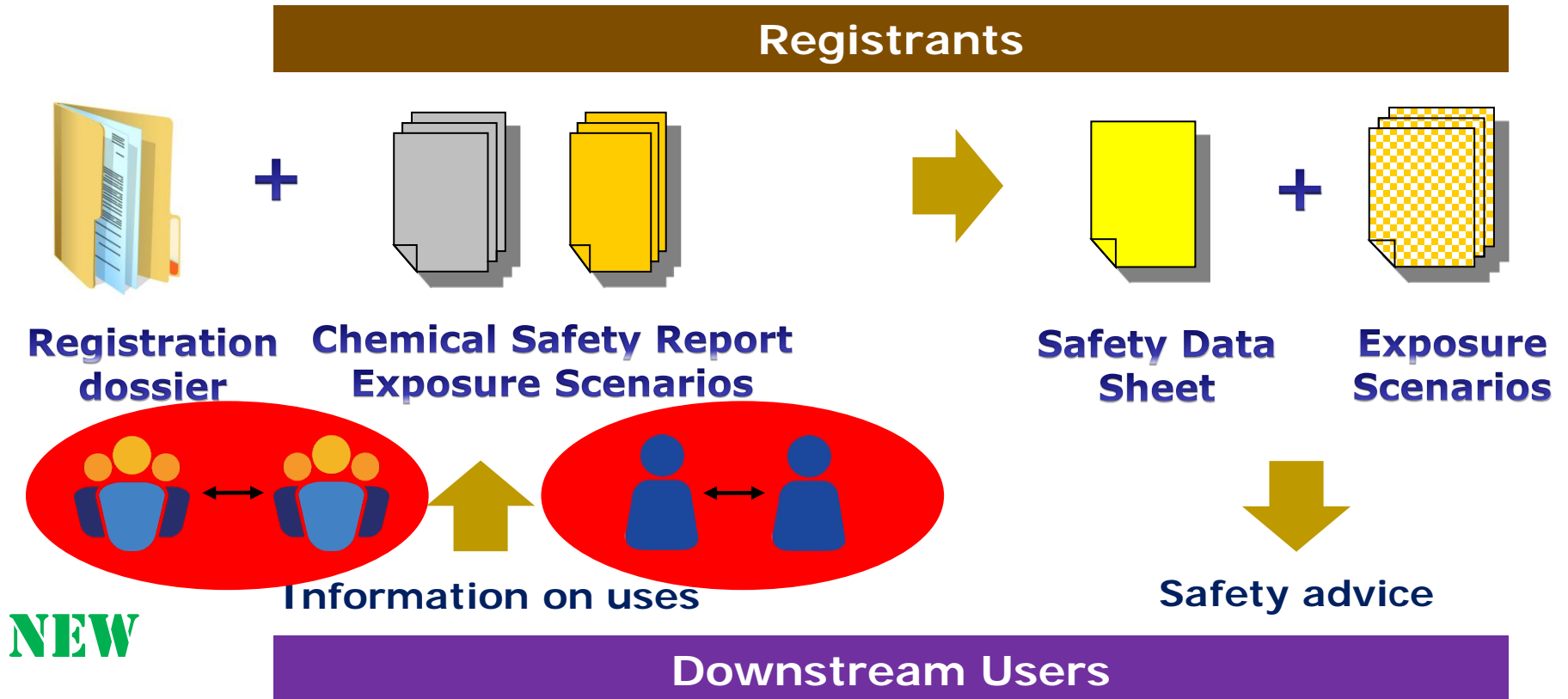


GES = Generic exposure scenario

SpERC = Specific environmental release category

SCED = Specific consumer exposure determinant

Organising communication in the supply chain



Communication in the supply chain

- **Where** to collect information
- **What** information to communicate
- In **which format** to communicate information
- **How** the supplier **can react** to the information passed on to him



Chapter 6: Communicating upstream



- Information on hazardous properties
- Information on inappropriate risk management measures
- Informing ECHA about new classification of a substance

Information on hazardous properties

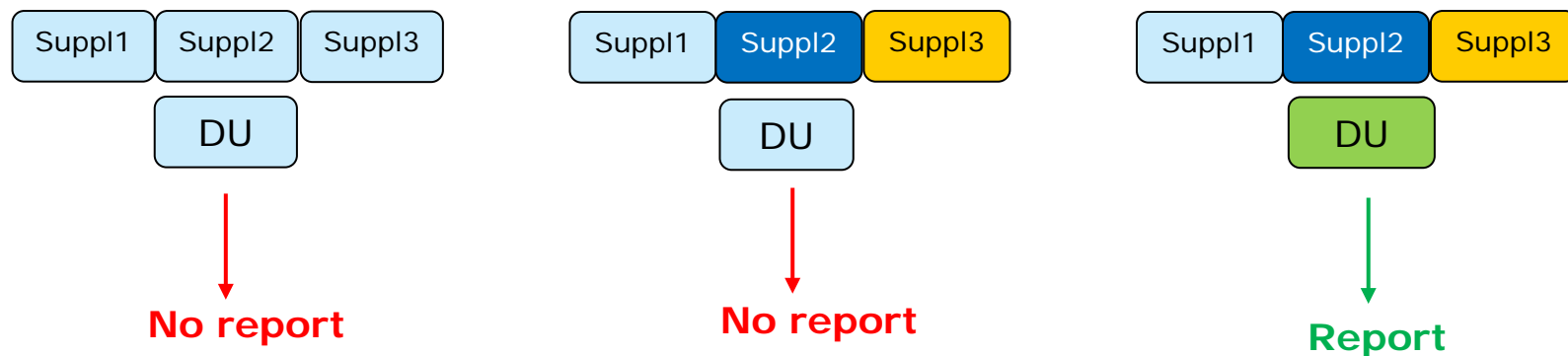
Inappropriate risk management measures

- How to decide whether information is **new**?
- What is "**inappropriate**" risk management measure?
 - E.g. not effective, overprotective
- How to **communicate**?
- What are the **timelines**?
- What is the **expected** supplier **response**?



New classification - Article 38(4) **NEW**

- When and what to report to ECHA



Note: Reporting classification differences is not required for a substance used in quantities less than 1 tonne per year

laura.walin@echa.europa.eu

www.echa.europa.eu

Subscribe to our news
echa.europa.eu/subscribe

Follow us on Twitter
[@EU_ECHA](https://twitter.com/EU_ECHA)