STUDY ON THE IMPACTS OF AUTHORISATION

WHAT DID WE LEARN IN ORDER TO IMPROVE THE STUDY IN 5 YEARS TIME?

Stock-taking conference on the implementation of REACH authorisation

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OUTLINE

What did we learn?

Which uncertainties might be resolved in 5 years time?

> How important are these data gaps and what can be done?

WHAT DID WE LEARN? - DATA SOURCES

Existing studies

Pre-survey engagement

Online surveys with NGOs and industry

Questionnaires to public authorities

Interviews

Market reports and statistics

WHAT DID WE LEARN? - DATA SOURCES

- > Existing studies Mostly qualitative information
 - > Not that many studies specific on REACH Authorisation
 - Some of the more recent authorisation studies were being carried in parallel difficult to incorporate their findings in a timely manner
 - > But it did provide a base for comparing our findings with
- > Pre-consultation (emails and calls) to raise awareness of the upcoming survey
 - Successful but very time consuming
 - Only a few NGOs have data on REACH authorisation
 - Lots of companies/sector groups were happy to speak but not many had necessary detailed data or were able/willing to share

Online survey

- Proved to be the approach that was able to provide get the most quantitative and monetary data
- Was a very long survey But in hindsight it should have been longer
- > It was very time consuming Should have been a separate study done in advance
- > NGO survey given small sample would have been better to do a questionnaire

eftec: Results of the study on the impacts of authorisation

WHAT DID WE LEARN? - DATA SOURCES

Questionnaires with public authorities

Successful as they are familiar with filling in questionnaires

Interviews

- Useful to provide further context/info to survey results
- Useful for producing case studies
- Did not reveal significantly more quantitative and monetary data
- Time consuming and difficult to arrange over summer period

Market reports and statistics

- Public statistics not always substance specific (e.g. groups of substances)
- > Market reports and statistics are not specific to uses subject to authorisation
- For substance specific case studies reports can be useful but not when trying to get a bigger picture on all SVHCs

WHAT DID WE LEARN - EASIER TO GET DATA AT SUBSTANCE LEVEL

1 EU LEVEL

Publicly available statistics

- Market reports

Interviews

- Online survey (dependent on sample size)

Existing studies

SUBSTANCE (SVHC) LEVEL

Publicly available statistics

✓ - Market reports

- Interviews / questionnaires

Online survey

- Existing studies

SPECIFIC USE OF AN SVHC



- Publicly available statistics
 - Market reports
 - Interviews / questionnaires
- Online survey
- Existing studies

LEVEL OF QUANTIFICATION POSSIBLE

EU market changes

Substitution

Costs

Benefits

Impacts on SMEs

WHAT UNCERTAINTIES MIGHT BE RESOLVED IN 5 YEARS TIME

- If some applicants were able to substitute or if they submit review reports
- Benefits of substitution (currently it is too early to say at the moment)

- Changes in exposure and emissions from continued uses of SVHCs (e.g. review reports)
- Any evidence of closure and relocation of production within the EU

Costs and benefits of enforcements of REACH authorisation

Comparison with findings from this study (i.e. some form of baseline has been established)

PRIORITISES FOR DATA GAPS

Changes in exposure of SVHCs

Changes in emissions of SVHCs

Benefits of substitution (A good or regrettable substitution?)

Evidence of closure and relocation and of diverted investment to non-EU sites

5 Competitiveness of EU companies

Comparison of the regulatory situation in other non-EU countries

ONE IDEA - A POST SUNSET DATE REPORT?

Report at substance level

Relevant for substances with lots of AfA's and/or high volumes being used

3 Applicants and alternative providers would have an incentive to input to it

Opportunity for NGOs to comment at substance level rather than per AfA

Information could also help applicants with their review reports / substitution efforts

May reduce concerns of granting longer review periods if EC get regular updates on the need for authorisation

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