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COMPLETE

Collector: Nano Consult - Non-Industry (Web Link)

Started:

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IP Address:

PAGE 2: Section I - Identification

Q1: Please provide the following details (*compulsory):

Your name:	MARCHE Patrice
Name of organisation* (if applicable):	INSERM
Town/City:	Grenoble
Country*:	France
E-mail address:	

Q2: Please indicate if you are responding to this questionnaire on behalf of/as:	c) a health and safety institute/academic organisation/research organisation
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Q3: Received contributions may be published on the Commission's website, with the identity of the contributor. Please state your preference with regard to the publication of your contribution:	My contribution may be published under the name indicated
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Q4: We might need to contact you to clarify some of your answers. Please state your preference below:	I am available to be contacted
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PAGE 3: Section III – Problem definition and objectives

Q5: Please rate the importance of the following objectives on a scale between 1 (not important at all) and 5 (very important).

- | | |
|--|---|
| a) Provide decision makers, regulatory authorities and professional users with information that allows for an appropriate response to health or environmental risks of nanomaterials | 5 |
| c) Maintain competitiveness and innovation of businesses bringing nanomaterials or products containing nanomaterials to the market (including SMEs) | 4 |
| d) Ensure consumer trust in products containing nanomaterials | 4 |
| e) Ensure the availability of relevant information on the presence of nanomaterials or products containing nanomaterials on the market | 3 |
| f) Ensure the proportionality of the information requirements and the associated costs and administrative burden. | 1 |
| g) Protect confidential business information | 2 |

Q6: To what degree (from 1 - not at all to 5 - fully) does the current legislative framework (including the REACH and CLP Regulations and product-specific legislation) and the currently available databases (including the JRC web platform, see http://ihcp.jrc.ec.europa.eu/our_databases/web-platform-on-nanomaterials) meet the following objectives?

- | | |
|--|---|
| a) Provide decision makers, regulatory authorities and professional users with information that allows for an appropriate response to health or environmental risks of nanomaterials | 4 |
| b) Provide consumers with relevant information on products containing nanomaterials on the market | 3 |
| c) Maintain competitiveness and innovation of businesses bringing nanomaterials or products containing nanomaterials to the market (including SMEs) | 3 |
| d) Ensure consumer trust in products containing nanomaterials | 2 |
| e) Ensure the availability of relevant information on the presence of nanomaterials or products containing nanomaterials on the market | 4 |
| f) Ensure the proportionality of the information requirements and the associated costs and administrative burden. | 2 |
| g) Protect confidential business information | 5 |

Q7: To what extent do you agree with the following statements from 1 (strongly disagree) to 5 (strongly agree):

- | | |
|---|---|
| a) The current level of available information on the presence of nanomaterials and products containing nanomaterials on the market is insufficient for an adequate response to health and environmental risks | 4 |
| b) The current level of available information on the presence of nanomaterials and products containing nanomaterials on the market is insufficient for informed consumer choice | 4 |
| c) The current level of available information on the presence of nanomaterials and products containing nanomaterials on the market is detrimental to consumer trust | 3 |
| d) The available information on the presence of nanomaterials and products containing nanomaterials on the market is presented in an incoherent or ineffective way | 4 |
| e) The establishment of national registries and notification schemes causes market fragmentation and hampers trade within the internal market | 2 |

<p>Q8: With regard to health and environmental hazards and risks of specific nanomaterials/types of nanomaterials, please tick the relevant boxes:</p>	<p>I am aware of health and/or environmental hazards of specific nanomaterials/types of nanomaterials</p> <p>,</p> <p>I am not aware of any classified nanomaterials,</p> <p>I am not aware of any DNELs/PNECs/OELs set for specific nanomaterials/types of nanomaterials</p> <p>,</p> <p>I am aware of significant exposure of workers/users/consumers to specific nanomaterials/types of nanomaterials</p>
<p>Q9: With regard to the past and current use of nanomaterials (tick the relevant box):</p>	<p>I am not aware of any health and/or environmental incidents which have occurred</p>
<p>Q10: The establishment of an EU nanomaterial registry (tick the relevant box):</p>	<p>Would significantly contribute to reducing the health and/or environmental risks related to the use of nanomaterials</p> <p>,</p> <p>If appropriate, please explain further: Nanomaterials are so diverse and difficult to precisely described, that registry would be of great value in providing references that can be shared.</p>

PAGE 5: Section V – Consumer trust

<p>Q11: In case information on the presence of nanomaterials in specific products were made available, what impact do you think this would have on consumers? (Please tick all that would apply)</p>	<p>c) Their purchasing decisions would not be affected</p>
<p>Q12: Do you believe that the public availability of information on the presence of nanomaterials in products would be likely to...(choose one of the following answers)</p>	<p>b) have no significant impact</p>

PAGE 6: Section VI - Innovation and competitiveness

<p>Q13: With regard to innovation, do you believe that information on nanomaterials and products containing nanomaterials that could be gathered in a nanomaterial registry would...(choose one of the following answers)</p>	<p>a) stimulate innovation (e.g. through increased consumer trust, increased awareness on nanomaterials)</p>
<p>Q14: With regard to competitiveness of EU companies manufacturing nanomaterials or products containing nanomaterials, do you believe that information on nanomaterials and products containing nanomaterials that could be gathered in a nanomaterial registry would...(tick all that apply)</p>	<p>a) stimulate intra-EU competitiveness,</p> <p>b) enhance the competitiveness of European companies against extra-EU companies</p>

PAGE 7: Section VIII – Possible options and exemptions

Q15: What would be the added value of a notification per use (i.e. for each mixture/article) compared to a notification per substance? – Please consider the usefulness of the information for public authorities, downstream user companies, workers and consumers.

The context of the uses of nanomaterial is as, if not more, important that their nature; how nanomaterial could disperse from its original packaging and enter in other environment including in body.

Q16: Which actors along the supply chain should be subject to notification requirements? (tick all that apply):

- a) Manufacturers of nanomaterials,
- c) Downstream users (e.g. re-formulators, manufacturers of products containing nanomaterials)
- ,
- d) Distributors to professional users (e.g. wholesalers)
- ,
- e) Distributors to consumers (e.g. retailers)

Q17: The following should be subject to notification requirements (tick all that apply):

- a) Substances,
- b) Mixtures containing nanomaterials,
- c) Articles with intended release of nanomaterials
- ,
- d) Articles containing nanomaterials without intended release

Q18: Is there a need to exempt certain types of nanomaterials?

No, all kinds of nanomaterials should be subject to notification obligations

Q19: Is there a need to exempt certain uses of nanomaterials?

No, all uses of nanomaterials should be subject to notification obligations

PAGE 8: Section IX – Nanomaterials Observatory

Q20: If a Nanomaterials Observatory is established instead of an EU-wide registry, what type of information should be collected? (please tick all that apply)

- a) Information from existing notification systems,
- b) Information from market studies on nanomaterials and products containing nanomaterials
- ,
- d) Information concerning products containing nanomaterials
- ,
- e) Information on the hazards and risks of nanomaterials

Q21: How should the information in a Nanomaterials Observatory be presented in order to reach the consumers, workers and authorities?

Recommendations associated to a label of quality, in the aim to propose guaranties which can be controlled.

Q22: In what ways could the information on nanomaterials from registries be potentially useful (tick all that apply):

- a) Risk assessment and/or risk management,
- b) Enforcement of worker protection,
- c) Promotion of safe use of nanomaterials in products
- ,
- d) Development of strategies to ensure the safe use of nanomaterials
- ,
- f) General education of the public

Q23: Please give a justification for your views (presented in the previous question) and describe which data would be necessary to allow the desired use (e.g. would information on substances alone be enough for informed consumer purchase decisions, or would this require information for each concerned product):

The nature of NM is not enough! Potential release, risk of exposure (inhalation, skin contact, ingestion by food of beverages ..) should be documented.

Q24: What would be the added value of a European nanomaterial registry beyond the current framework of chemicals legislation, including REACH registration?

A label of quality and independent controls. Controls have to be done by entities independent from producers, developers, purchasers to be reliable and trusted by public. consumers organizations could be part of the control process.

Q25: Please provide any other comments that you would like to share regarding transparency measures for nanomaterials on the market.

It is important to set the quantity of information to be delivered to the public. An excess of information may be deleterious for a wise opinion.