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COMPLETE

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PAGE 2: Section I - Identification

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

Your name:	Jeanette
Country*:	Sweden

Q2: Please indicate if you are responding to this questionnaire on behalf of/as:	a) an individual
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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
b) Provide consumers with relevant information on products containing nanomaterials on the market	5
c) Maintain competitiveness and innovation of businesses bringing nanomaterials or products containing nanomaterials to the market (including SMEs)	1
d) Ensure consumer trust in products containing nanomaterials	1
e) Ensure the availability of relevant information on the presence of nanomaterials or products containing nanomaterials on the market	5
f) Ensure the proportionality of the information requirements and the associated costs and administrative burden.	1
g) Protect confidential business information	1
Please provide additional comments	1. There are serious issues with nanomaterials, and I don't want them in anything. 2. Keep the corporate lobbyists far away from the parliamentarians and commissioners, and make sure that EFSA is completely free from corporate influence and corruption.

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	1
b) Provide consumers with relevant information on products containing nanomaterials on the market	1
c) Maintain competitiveness and innovation of businesses bringing nanomaterials or products containing nanomaterials to the market (including SMEs)	5
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	1
f) Ensure the proportionality of the information requirements and the associated costs and administrative burden.	Do not know
g) Protect confidential business information	Do not know

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	5
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	5
c) The current level of available information on the presence of nanomaterials and products containing nanomaterials on the market is detrimental to consumer trust	1
d) The available information on the presence of nanomaterials and products containing nanomaterials on the market is presented in an incoherent or ineffective way	5
e) The establishment of national registries and notification schemes causes market fragmentation and hampers trade within the internal market	1

Q8: With regard to health and environmental hazards and risks of specific nanomaterials/types of nanomaterials, please tick the relevant boxes:	<div>I am aware of health and/or environmental hazards of specific nanomaterials/types of nanomaterials</div> <div>,</div> <div>I am aware of specific nanomaterials that are classified as hazardous under Regulation (EC) No 1272/2008 on classification, labelling and packaging of substances and mixtures</div> <div>,</div> <div>I am aware of significant exposure of workers/users/consumers to specific nanomaterials/types of nanomaterials</div>
Q9: With regard to the past and current use of nanomaterials (tick the relevant box):	<div>I am aware of health and/or environmental incidents which have occurred</div>
Q10: The establishment of an EU nanomaterial registry (tick the relevant box):	<div>Would significantly contribute to reducing the health and/or environmental risks related to the use of nanomaterials</div> <div>,</div> <div>If appropriate, please explain further: A nanomaterial registry is useless if no information about risks are added.</div>

PAGE 5: Section V – Consumer trust

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)	<div>b) They would try to avoid those products,</div> <div>d) They would search for more information</div>
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)	<div>b) have no significant impact,</div> <div>Comments: What type of information are you referring to? Who’s going to provide the (unbiased) information? I have to tick (b) here since the question is impossible to answer.</div>

PAGE 6: Section VI - Innovation and competitiveness

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)	<div>b) have no significant impact on innovation</div>
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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)

- e) hamper intra-EU competitiveness,
- f) hamper the competitiveness of European companies against extra-EU companies
- ,
- Please explain
- I think the EU-28 should be a forerunner regarding clean, risk-free and healthy products. It's insane to do otherwise.

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.

I honestly don't know.

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

- a) Manufacturers of nanomaterials,
- b) Importers 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?

Respondent skipped this question

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
- ,
- c) Information on the use of nanomaterials across Europe
- ,
- d) Information concerning products containing nanomaterials
- ,
- e) Information on the hazards and risks of nanomaterials
- ,
- f) Other (please explain):
And again . . . we want unbiased information.

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

On the appropriate Public authorities webpages, for example the Public Health Agency, National Food Agency etc.

I also see upcoming problems regarding health insurance.

PAGE 9: Section X - Potential use and benefits of a nanomaterial registry

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,
- e) Informed purchasing decisions by consumers,
- f) General education of the public,
- g) Other purposes (please specify)
- c) maybe d) hardly

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):

I think a combination of both.

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

If presented in a understandable manner it would add tremendous value.

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

I'm sorry, but I do smile when I read the word "transparency". The European institutions really suffers from a grave transparency deficit.