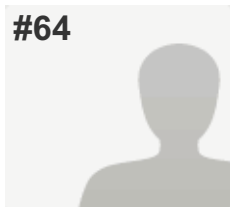


#64



**COMPLETE**

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

**Started:**

**Last Modified:**

**Time Spent:**

**IP Address:**

## PAGE 2: Section I - Identification

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

Organisation\*:

Country\*:

UK

Contact name:

E-mail address:

**Q2: 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 but should be kept anonymous

**Q3: We might need to contact you to clarify some of your answers. Please state your preference below:**

I am available to be contacted

**Q4: Did your organisation participate in the online survey (undertaken by RPA/BiPRO for the European Commission in early 2014) on the administrative burden of the notification schemes?**

Do not know

## PAGE 3: Section II - Organisation Information

**Q5: Please indicate which of the following applies to you or your members (tick all that apply):**

d) is an importer of nanomaterials

**Q6: Please indicate the four-digit NACE code of your primary and secondary business sector (if applicable). If you require information regarding NACE codes, please visit the European Commission Competition webpage at [http://ec.europa.eu/competition/mergers/cases/index/nace\\_all.html](http://ec.europa.eu/competition/mergers/cases/index/nace_all.html)**

Primary business sector (NACE 4 digit code):

C26.7

**Q7: Please indicate the number of employees.**

≥ 250 employees

**Q8: Please indicate the approximate annual turnover of your organisation and the annual turnover which relates to nano-related products (where these include nanomaterials as well as mixtures and articles containing nanomaterials).**

Annual turnover

≤ €250k

Nano-related annual turnover

≤ €250k

**Q9: Please indicate the number of nano-related products (where these include nanomaterials as well as mixtures and articles containing nanomaterials) that you place on the national market.**

Nanomaterials	less than 6
Mixtures	less than 6
Articles	less than 6

**Q10: Please indicate the number of nano-related products (where these include nanomaterials as well as mixtures and articles containing nanomaterials) that you place on the EU market.**

Nanomaterials	less than 6
Mixtures	less than 6
Articles	less than 6

**Q11: Please indicate the number of nano-related products (where these include nanomaterials as well as mixtures and articles containing nanomaterials) that you place on the global market.**

Nanomaterials	less than 6
Mixtures	less than 6
Articles	less than 6

**Q12: Please indicate the number of customers and, if applicable, number of suppliers for all your nano-related products combined (where these include nanomaterials as well as mixtures and articles containing nanomaterials).**

Number of customers	less than 6
Number of suppliers	less than 6

**Q13: 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	4
b) Provide consumers with relevant information on products containing nanomaterials on the market	2
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	3
e) Ensure the availability of relevant information on the presence of nanomaterials or products containing nanomaterials on the market	2
f) Ensure the proportionality of the information requirements and the associated costs and administrative burden.	3
g) Protect confidential business information	5

**Q14: 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](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	2
b) Provide consumers with relevant information on products containing nanomaterials on the market	2
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	3
f) Ensure the proportionality of the information requirements and the associated costs and administrative burden.	4
g) Protect confidential business information	4

**Q15: 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   | 4 |
| d) The available information on the presence of nanomaterials and products containing nanomaterials on the market is presented in an incoherent or ineffective way  | 3 |
| e) The establishment of national registries and notification schemes causes market fragmentation and hampers trade within the internal market   | 3 |

**PAGE 5: Section IV – Health and environmental aspects**

**Q16: With regard to health and environmental hazards and risks of specific nanomaterials/types of nanomaterials, please tick the relevant boxes:**

I am aware of health and/or environmental hazards of specific nanomaterials/types of nanomaterials

,

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

**Q17: With regard to the past and current use of nanomaterials (tick the relevant box):**

I am aware of health and/or environmental incidents which have occurred

**Q18: The establishment of an EU nanomaterial registry (tick the relevant box):**

I do not know

**PAGE 6: Section V – Consumer trust**

**Q19: In case information on the presence of nanomaterials in your products were made available, what impact do you think this would have on your clients? (Please tick all that would apply)**

d) They would search for more information

**Q20: 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)**

b) have no significant impact

PAGE 7: Section VI - Innovation and competitiveness

<b>Q21: 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)</b>	a) stimulate innovation (e.g. through increased consumer trust, increased awareness on nanomaterials)
<b>Q22: 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)</b>	a) stimulate intra-EU competitiveness, b) enhance the competitiveness of European companies against extra-EU companies

PAGE 8: Section VII – Possible impact of a registry on your company/members of your association

<b>Q23: Overall, how would a possible obligation to notify nanomaterials at the EU level affect your company/the members of your association, assuming that no exemptions were to be made from 1 (no impact) to 5 (significant impact):</b>	
a) with respect to nanomaterials on their own	2
b) with respect to nanomaterials in mixtures	3
c) with respect to articles with intended release of the nanomaterials	3
d) with respect to articles containing nanomaterials in general (i.e. in case also articles without an intended release of nanomaterials were to be covered)	2
<b>Q24: Would disclosure of the notified information conflict with the confidentiality of business information?</b>	Yes, there would be a conflict with business information confidentiality
<b>Q25: Do you experience or expect any significant barriers for your company/members of your association from diverging registration obligations in the schemes in France/Belgium/Denmark?</b>	No, we do not expect any barriers
<b>Q26: Is the market for your nanomaterials/products containing nanomaterials significantly different from Member State to Member State?</b>	Yes, the markets differ at national level
<b>Q27: In case the European Commission were to recommend a best practice model for national notification schemes based on the experiences in France, Belgium and Denmark, which elements of these systems can be considered as “best practice”?</b>	<i>Respondent skipped this question</i>

PAGE 9: Section VIII – Possible options and exemptions

**Q28: 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.**

Regular testing, Quality assurance, more accurate reporting. This could potentially be downgraded at a future point.

**Q29: 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)
- ,

Please explain:

Once formulated the product is known and understood, prior to this downstream users may well change the properties. Manufacturers and importers will give wider breadth to the reporting - more of a catch all.

**Q30: The following should be subject to notification requirements (tick all that apply):**

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

**Q31: Is there a need to exempt certain types of nanomaterials?**

Yes, certain types of nanomaterials should be exempted from a notification system

,

If yes, which types should be exempted and why? (in terms of specific properties, available knowledge, absence of hazards, etc.)  
I would think if there was evidence of significant use of a nanomaterial over a long period, this could be exempted. For example casein micelles would make milk a nanomaterial - do we really need this reported?

**Q32: Is there a need to exempt certain uses of nanomaterials?**

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

**Q33: 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

**Q34: How should the information in a Nanomaterials Observatory be presented in order to reach the consumers, workers and authorities?**

I suspect it would be a single website with consolidated information that would need a high level of ongoing expert maintenance.

**PAGE 11: Section X - Potential use and benefits of a nanomaterial registry**

**Q35: 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
- ,
- e) Informed purchasing decisions by consumers,
- f) General education of the public

**Q36: 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):**

*Respondent skipped this question*

**Q37: What would be the added value of a European nanomaterial registry beyond the current framework of chemicals legislation, including REACH registration?**

Current legislation does not adequately consider that a nanomaterial has properties that are not currently addressable within the existing frameworks. Given that nanomaterials in many areas have been shown to have significantly different properties, this concern should not be ignored.

**Q38: Please provide any other comments that you would like to share regarding transparency measures for nanomaterials on the market.**

*Respondent skipped this question*