

Public consultation on alternatives

Seminar on Applications for Authorisation

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Outline

- Why do we need a Public Consultation on alternatives?
- How does it work?
- What information on alternatives is requested?
- How is the submitted information evaluated?
- How to ensure a meaningful and efficient Public Consultation on Alternatives?

Statistics on received comments

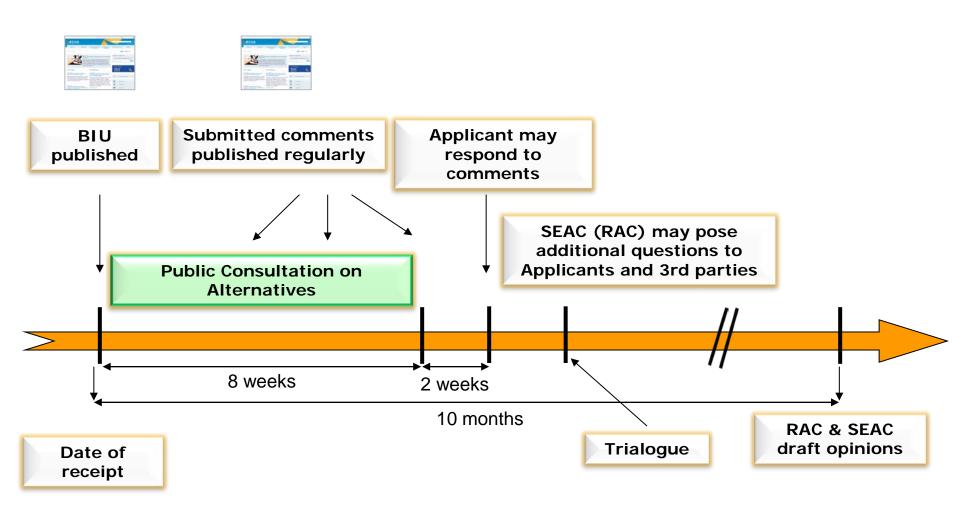


Purpose

- To gather information on alternatives
 - Relevant and meaningful information on alternatives is gathered in support to the Committees' scrutiny
- Public engagement in the regulatory process
 - All available information on alternatives is taken into account in the decision for granting an authorisation
 - Transparency and trustworthiness



Process overview





Process

- Start of the Public Consultation:
 - upon receipt of application (i.e. fee paid by the applicant)

publication of the Broad information on uses package

on ECHA's website:

Mid-February

Mid-May

Mid-August

Mid-November



http://www.echa.europa.eu/addressing-chemicals-of-concern/authorisation/applications-for-authorisation





ECHA > Addressing Chemicals of Concern > Authorisation > Applications for authorisation

Applications for authorisation

The placing on the market and use of Substances of Very High Concern included in the Authorisation List requires an authorisation. A manufacturer, an importer or a downstream user can apply for an authorisation. Applications for authorisation are submitted to ECHA. At the end of the authorisation process, which includes a public consultation and the development of opinions by ECHA's Committees on Risk Assessment and Socio-economic Analysis, the European Commission decides on the granting or refusing of authorisations.

Public consultation

The application for authorisation process includes a period of public consultation.

Anyone can comment on the uses of the substance related to the application, in particular to provide information on alternative substances or technologies. Those most likely to be interested are companies, organisations representing industry or civil society, individual citizens, as well as public authorities.

The public consultation lasts for eight weeks.

Provide your comments

In order to facilitate the work of ECHA's Committees in reviewing the comments received, you are kindly invited to provide your comments in English preferably. By clicking on a link in the table below, you will get access to the full broad information on the use applied for, and to the related commenting form.

Comments are welcomed from the EU or beyond.

Consultation O	Substance C Name	EC Number ©	CAS Number ©	Deadline for comments	Applicant (s)	Broad information on cuse applied for		
001-01	Substance Test 1	300-000-0	300-00-00		Applicants Test 1	BIU Test 1	Details	K
002-01	Substance Test 2	400-000-0	400-00-00		Applicants Test 1	BIU Test 1	Details	



Information on use applied for in Applications for Authorisation «Back to Consultation list

Substance Name	Substance Test 1	
EC Number	300-000-0	
CAS Number	300-00-00	
Annex XIV identifier		
Broad information on use applied for	BIU Test 1	Finalised by ECHA based on
Broad information on use applied for (name and conditions of use)		applicant's proposal
Use applied for number in application for authorisation		
Broad information on use applied for (Use descriptor system)		
Exposure scenario (non confidential)		Published as provided by
Analysis of Alternatives (non confidential report)		the applicant. Applicant's
Substitution Plan (non confidential summary)		responsibility to ensure no confidential info is included
Socio Economic Analysis (non confidential summary)		confidential into is included
Consultation Number	001-01	
Applicant(s)	Applicants Test 1	8 weeks of Public
Application type		Consultation on Alternatives
Start of consultation		
Deadline for comments		Submitted comments
Comments	Give comments	published on website



Comments on public consultation 0001-01

Summary

The following comments have been submitted to date as part of the public consultation on alternatives for Applications for authorisation. ECHA accepts no responsibility or liability with regard to the information (including attachments) presented below.

For details on the public consultation process, please refer to Applications for Authorisation.

Substance name Bis(2-ethylhexyl) phthalate (DEHP)

EC Number 204-211-0
CAS Number 117-81-7

Entry Nr in Annex XIV

Broad information on use applied for (title) The processing of a stop-off formulation containing DEHP during the diffusion bonding and

manufacture of aero engine fan blades.

Consultation number 0001-01

Applicant name(s) Rolls-Royce plc

Consultation period 14/08/2013 - 09/10/2013

Response to comments by applicant

New comments added regularly

Comments

#	Submitter	Alternative						
		Туре	Generic name	EC Number	CAS Number	Description of technology	Classification and Labelling	Attachments
9	Affiliation: organisation	Other				DEHP elimination		Download
	Type/ Role in supply chain: Non-governmental organisation (NGO) Name of org/company:							
	European Environmental Bureau (EEB)							
	Country: Belgium							



Requested information on alternatives during Public Consultation

- Substance/mixture identification and/or description of technology/process
- Substance properties
- Classification & labelling information
- Technical feasibility
- Economic feasibility
- Risks
- Availability

The web form allows parties to submit the same comment for several ongoing consultations



How RAC and SEAC evaluate the submitted information?

- Assess submissions (together with the application) based on:
 - Relevance for the uses applied for
 - Quality and clarity
 - Completeness of the analysis: technical feasibility, economic and risk information

• Trialogue:

- How relevant are the public consultation submissions to the uses applied for?
- Opportunity to clarify questions on alternatives with the applicant and, if required, third parties



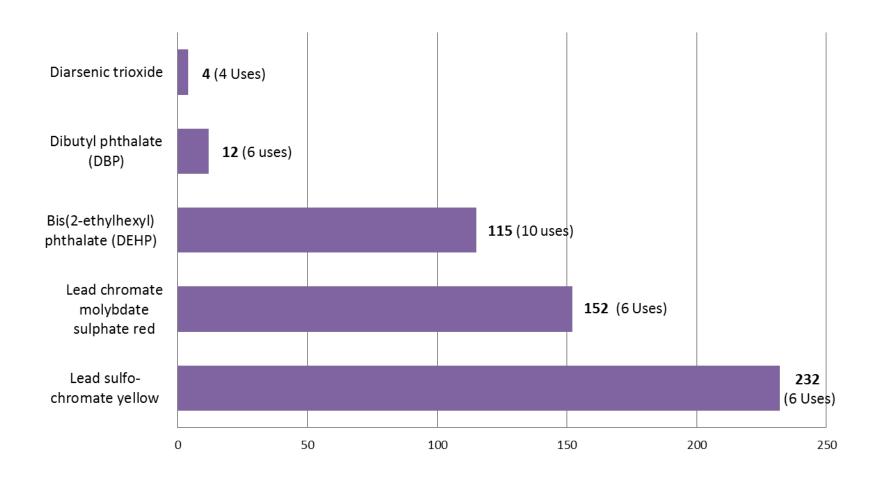
BIU package: key to meaningful consultation

Communicate well your case to the public to maximise relevant submissions from 3rd parties & to limit the number of follow up questions from SEAC/RAC

- Trade off between releasing information in the public domain (as part of the BIU package) and having to respond to irrelevant information submitted during the Consultation
 - Can you communicate your case using publicly available information?
 (e.g., using ranges, industrial averages, patent information, etc.)
- Anticipate what can be submitted during the Public Consultation
 - Screen publicly available information on alternatives
 - "Test drive" your BIU package to determine what alternatives may come up based on your public submission only

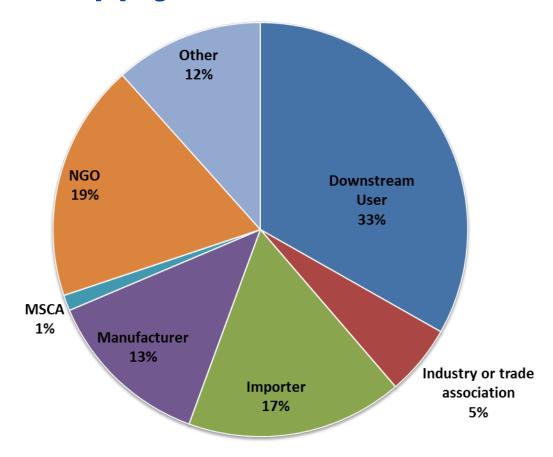


Received comments per substance (Uses)





Type of organisation or company and its role in the supply chain





Key messages

- Public Consultation:
 - identification of information gaps on alternatives for the use applied for
 - third parties involvement and support to the Committees' scrutiny
- BIU package: key to meaningful and efficient consultation on alternatives
- Possibility for applicant to provide counterarguments but with tight deadlines
 - Responses to submissions within 2 weeks of close of consultation
 - Trialogue
 - Responses to RAC/SEAC written questions

Thank You!

