

Introduction

Webinar: New ECHA public data
availability system – Part II

19 October 2023

Roberta DI BLASI
European Chemicals Agency



What you can expect today

Time	Topic	Speaker
11.00	Introduction	Roberta DI BLASI
11.05	Recap with timeline	Roberta DI BLASI
11.10	Live demo (REACH registrations)	Roberta DI BLASI
11.20	Regulatory obligations and processes	Gerson MARTIN
11.25	Transition plan and conclusions	Gerson MARTIN
11.00 – 12.00	Slido.com open for questions	

Live Q&A

- Join Q&A at: [slido.com](https://www.slido.com)
Event code:
- Send questions from
11:00 to 12:00 (EEST, GMT +3), October
19, 2023
- Only questions within scope
- Question not answered?
Contact us: echa.europa.eu/contact

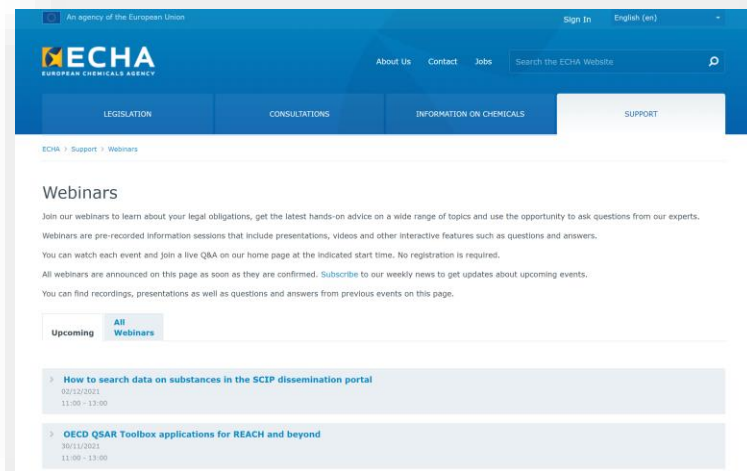


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Material available

- Video recording
- Presentations
- Q&A transcript (soon after the event)

- Subscribe to our newsletter at echa.europa.eu/subscribe



The screenshot shows the ECHA website's 'Webinars' page. The header includes the ECHA logo, navigation links (About Us, Contact, Jobs), a search bar, and a menu with categories: LEGISLATION, CONSULTATIONS, INFORMATION ON CHEMICALS, and SUPPORT. The main content area is titled 'Webinars' and contains the following text:

Join our webinars to learn about your legal obligations, get the latest hands-on advice on a wide range of topics and use the opportunity to ask questions from our experts. Webinars are pre-recorded information sessions that include presentations, videos and other interactive features such as questions and answers. You can watch each event and join a live Q&A on our home page at the indicated start time. No registration is required. All webinars are announced on this page as soon as they are confirmed. [Subscribe](#) to our weekly news to get updates about upcoming events. You can find recordings, presentations as well as questions and answers from previous events on this page.

Below the text, there is a filter for 'Upcoming' and 'All Webinars'. Two webinar events are listed:

- > **How to search data on substances in the SCIP dissemination portal**
02/12/2021
11:00 - 13:00
- > **OECD QSAR Toolbox applications for REACH and beyond**
06/11/2021
11:00 - 13:00

echa.europa.eu/webinars

Recap with updated timeline

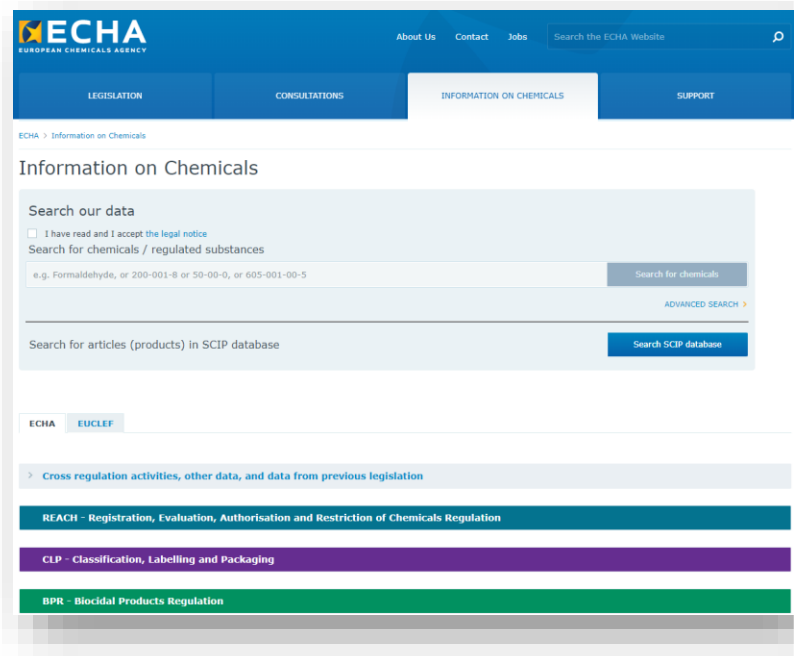
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European Chemicals Agency



ECHA's public chemicals database



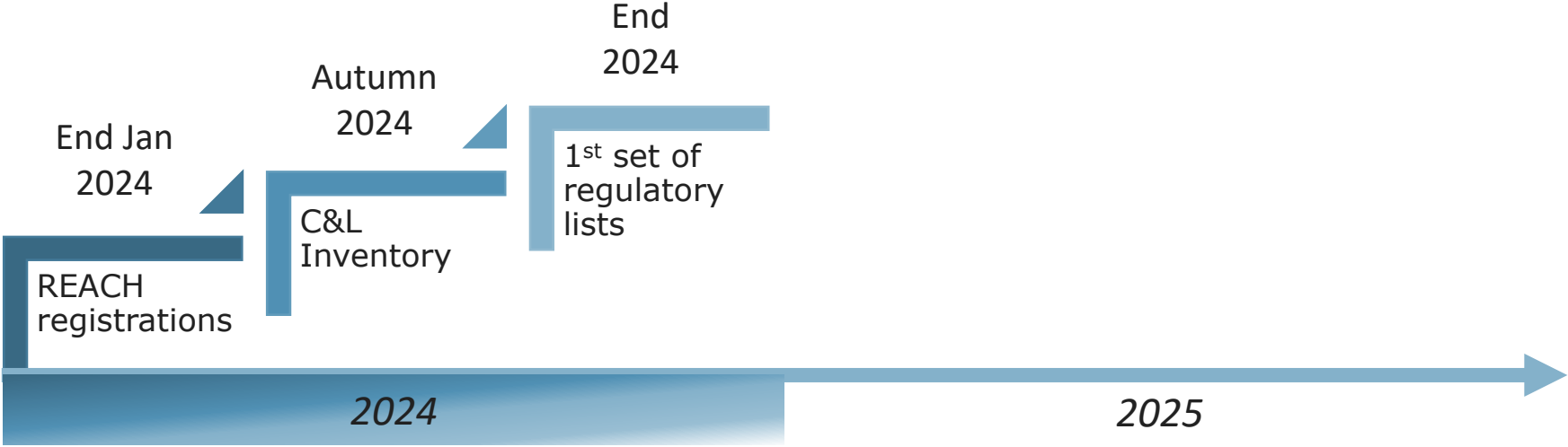
Getting ready for the future

New legislation and tasks
Increased quantity and wider
variety of data

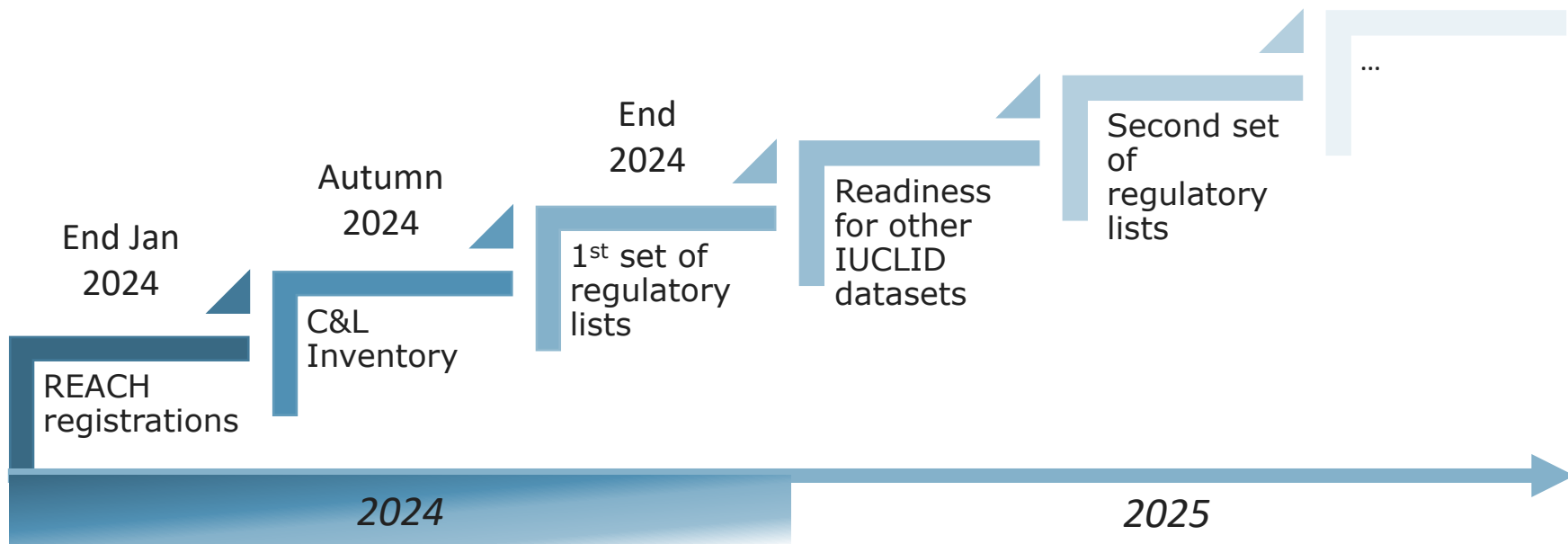
Key aims

Modular, expandable, scalable
Improved performance
Enhanced user experience

Building the new system



Building the new system



Summary

- Development of first version of the new data availability system is advancing well
- Go live end of January 2024 with REACH registrations
- Development by adding new data sets and enhancing the system features
- Stakeholder feedback very useful. Will continue user engagement also after go-live to collect feedback



Live demo

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Thank you

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Regulatory obligations and processes

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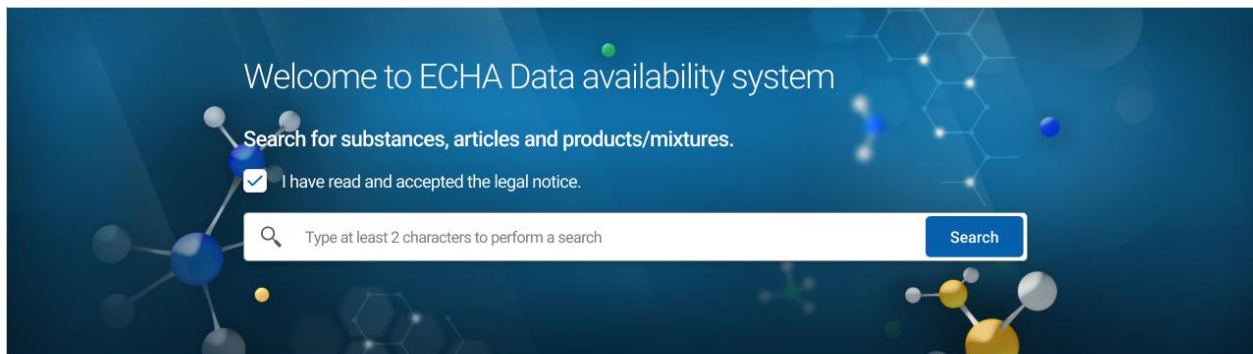
Gerson MARTIN
European Chemicals Agency



Regulatory obligations and processes

- **Regulatory obligation lists** – represent the current legal status of the entries
- **Regulatory processes** - provide insight on which substances are subject to regulatory activities and at which stage of the process they are
- Incrementally integrated into the new data availability system

Landing page



Welcome to ECHA Data availability system

Search for substances, articles and products/mixtures.

I have read and accepted the legal notice.


[Search](#)

Search engine

Regulatory data

Regulatory obligations  [View all obligations >](#)

- Candidate list [REACH](#)
- Active substance approval [BPR](#)
- Restriction list [REACH](#)
- Harmonised C&Ls [CLP](#)
- Authorisation list [REACH](#)

Regulatory processes  [View all processes >](#)

- Registry of intentions - CLH [CLP](#)
- Dossier evaluation status [REACH](#)
- CoRAP list (substance evaluation) [REACH](#)
- ED assessment list [Cross-regulation](#)
- Registry of intentions - SVHC [REACH](#)

Direct access to lists and processes

View all regulatory obligation lists

Filters:

9 Regulatory obligations, Rows per page

Regulatory obligation	Relevant legislation	Organisation
Authorization list REACH Annex XIV. Substances for which authorisation is required to be allowed to use the substance.	REACH	ECHA
Biocidal products authorised on the EU/EEA market	BPR	ECHA
Candidate list of SVHC for authorization Substances to be considered for potential authorisation at EU level.	REACH	ECHA
Chemicals subject to PIC	PIC	ECHA
Inland Transport of Dangerous Goods Directive, Annex III, ADN	Inland Transport of Dangerous Goods	DG MOVE
List of authorised biocidal products Biocidal products authorised to be on the market in the EU.	BPR	ECHA
List of harmonised classifications (Annex VI)	CLP	ECHA

Features

- Sorting
- Filtering
- Organisation

Regulatory obligation list

← Substances restricted under REACH ?

Intentions and Annex XV restriction proposals received by ECHA

REACH

The table below is the Annex XVII to REACH and includes all the restrictions adopted in the framework of REACH and the previous legislation, Directive 76/769/EEC. Each entry shows a substance or a group of substances or a substance in a mixture, and the consequent restriction conditions. The latest consolidated version of REACH...

[View more](#) ▾

Filter by substance identifiers



Search for substance identifiers e.g. Formaldehyde, or 200-001-8 or 50-00-0, or 605-001-00-5

Search





Additional Filters ▾

Entities ?

Show indirect entries as well: No

50 Entities, Rows per page

Showing 1-50 of 50 < Prev | Next >

Substance Identifiers			Process attributes			
Substance name ⌵	EC / List No. ⌵	CAS No. ⌵	Entry no. ⌵	Conditions ⌵	Appendices ⌵	Details ⌵
4,4'-isopropylidenediphenol 	262-968-2	61788-33-8	01	 01.pdf	Appendix 7	Restriction details
Lorem Ipsum 	262-968-2	61788-33-8	02	 02.pdf	Appendix 7	Restriction details

Short description

Filtering

Link to details page

Regulatory obligation entry details

Restriction Details ?

Intentions and Annex XV restriction proposals received by ECHA

REACH

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[View Restriction list](#)

Entry information

Entry no.	01
Q&As	<ul style="list-style-type: none">10781198
Remarks	Lorem ipsum dolor sit amet, consectetur adipiscing elit, sed do eiusmod tempor incididunt ut labore et dolore magna aliqua.
Restriction conclusion	Lorem ipsum dolor sit amet, consectetur adipiscing elit, sed do eiusmod tempor incididunt ut labore et dolore magna aliqua.

Timeline Expand all / Collapse all

- Restriction list inclusion 25-09-2007**
Sep 2007
[View more info](#) v
- Restriction list inclusion 22-06-2009**
Jun 2009
[View more info](#) v
- Restriction list inclusion 25-05-2023**
May 2023

Short description

General info about the entry

Timeline of events

Regulatory processes

The screenshot displays the ECHA platform interface for 'Regulatory processes'. At the top left is the ECHA logo and 'platform logo'. A 'Log in' button is at the top right. The main heading is '← Regulatory processes' with a help icon. Below this is a filter section with a search box 'Search for Regulatory process', a 'Select Regulation' dropdown, and a 'Select Organisation' dropdown. The main content area shows '11 Regulatory obligations, Rows per page 100' and 'Showing 1-11 of 11' with navigation arrows. A table lists regulatory processes with columns for 'Regulatory process', 'Relevant legislation', and 'Organisation'. The table contains 11 rows of data.

Regulatory process	Relevant legislation	Organisation
Assessment of regulatory needs (ARN)	Lorem ipsum	ECHA
Biocidal active substance approval	BPR	ECHA
Dossier evaluation status Details of dossier compliance checks and testing proposal evaluation.	REACH	ECHA
Endocrine disruptor assessment list	Lorem ipsum	ECHA
List of substances proposed as POPs	POP	ECHA
PBT assessment list Substances undergoing a PBT/vPvB assessment.	Lorem ipsum	ECHA
Recommendations for inclusion in the Authorisation List Substances included in ECHA's draft or final recommendations for inclusion in Annex XIV (Authorisation List)	REACH	ECHA

Features

- Sorting
- Filtering
- Organisation

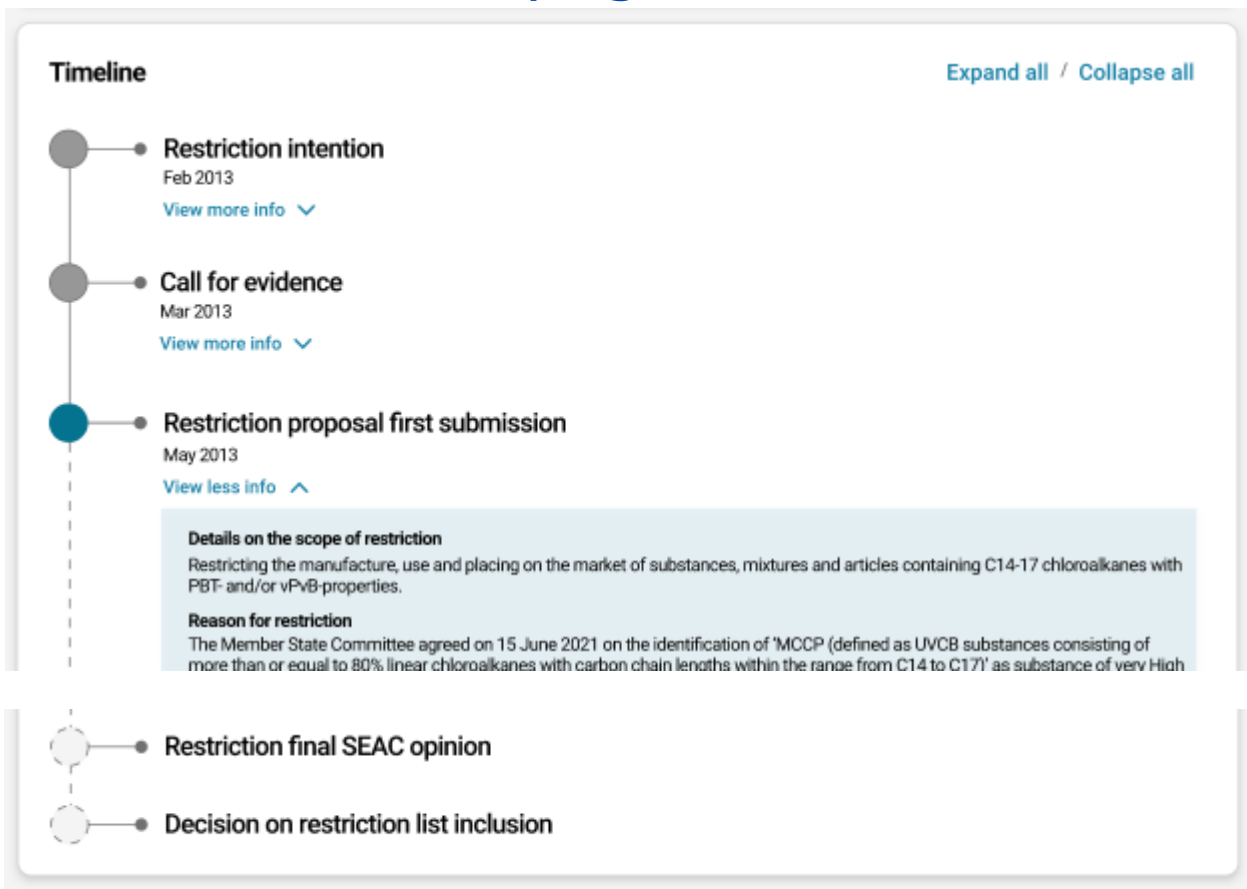
Process details page

The screenshot shows the ECHA platform interface for the Ethanol process. The top navigation bar includes the ECHA logo, a 'platform logo' button, and a 'Log in' button. Below this, the process name 'Ethanol' is displayed with a back arrow and a flask icon. Regulatory information is provided: 'Regulatory process name: ETH', 'Lorem ipsum: AGVJD9', 'EC number: 200-578-6', and 'CAS number: 64-17-5'. A left-hand sidebar contains navigation links: Overview, Identity, Relations, Dossiers, Classification & Labelling, and Regulatory context (with sub-links for Obligations and Processes). The main content area is titled 'Registry of restriction intentions until outcome' with a back arrow, a help icon, and a 'REACH' tag. It includes a sub-header 'Intentions and Annex XV restriction proposals received by ECHA' and a paragraph explaining the registry's purpose. A 'View more' link is present. Below this is a 'Case summary' box with the following details:

Case status	ONGOING
Submitter	France
Current conclusion on the Restriction	-
Current stage	Restriction proposal first submission
Current stage date	12 May 2013
Details on the scope of restriction	Restricting the manufacture, use and placing on the market of substances, mixtures and articles containing C14-17 chloroalkanes with PBT- and/or vPvB-properties.
Reason for restriction	The Member State Committee agreed on 15 June 2021 on the identification of 'MCCP (defined as UVCB substances consisting of more than or equal to 80% linear chloroalkanes with carbon chain length with the range from C14 to C17) substances of very high concern (SVHC) under the REACH Regulation.

Summary of the process information

Process details page



Timeline of events

- Grey: past events
- Blue: latest event
- Dashed: future steps

Transition plan and conclusions

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Transition plan

- Incremental approach



- Continuous access to information



- We will continue engaging with you



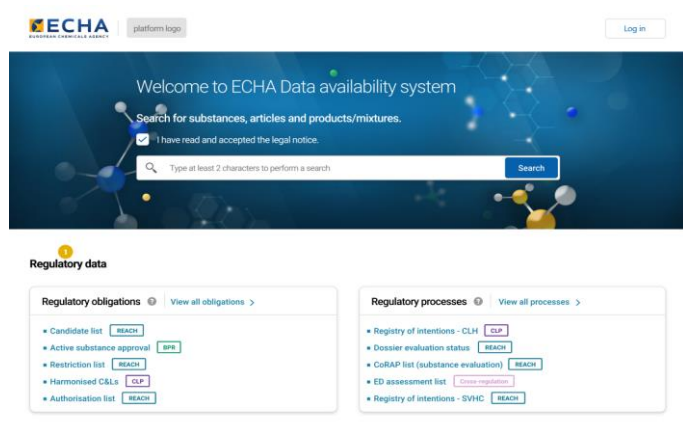
- Support material will be prepared to guide you in this journey

- E.g., new dedicated page on ECHA's website to inform you about progress (January 2024)
- Webinar series will continue

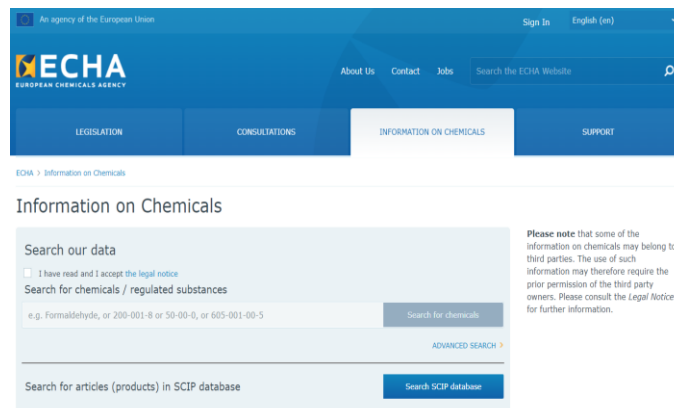


Transition plan

- Go live end of January 2024
- Two parallel systems



- REACH registrations
 - + C&L Inventory ~ Autumn 2024
 - + Regulatory lists ~ End 2024



Everything else

Live Q&A

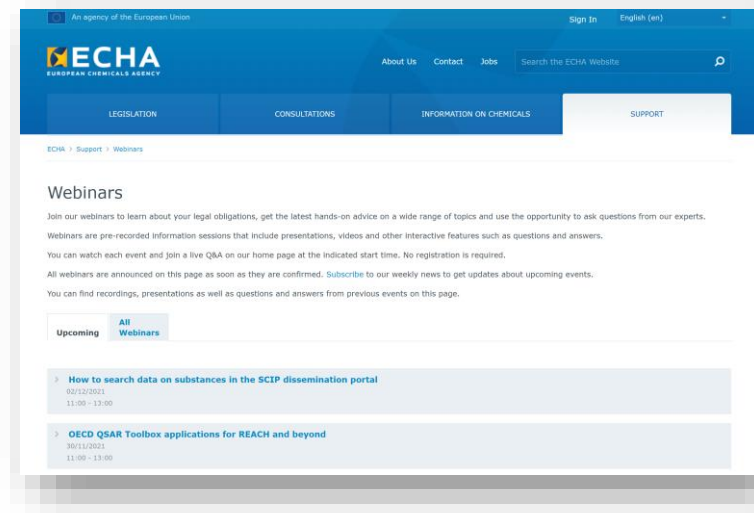
- Join Q&A at: slido.com
Event code:
- Webinar open until 12:00 Helsinki time (EEST, GMT+3) to answer questions
- If your question is not answered by the end of the webinar, send it via our contact form: echa.europa.eu/contact



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- Presentations
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