Submission for Restrictions

A submission via the corresponding webform is required for any initiation or update of an activit undertaken by an authority. This allows ECHA to manage the activity in an efficient manner and to inform other authorities and stakeholders via the Activity Coordination Table (ACT) or its public version (PACT) witl up to date information on the activity undertaken.

Fields marked with * are required. Other information is only required if changed from previous submissions.

Submission related information

Please specify below the type of submission: "Intention" announces the start of the activity, "Submission o documentation" refers to any document or dossier submitted during the process (e.g. outcome document Annex XV dossier, RCOM) and "Update of information" refers to any other change in the activity. If this is "Withdrawal" of the activity, please provide reasoning under "Other remarks" or as an attachment.

Type of submission : *

Intention

Activity performed by

Specify the organisation who is responsible for performing this activity. In case, activities on this substanc are performed by more than one Member State Authority (e.g. Substance Evaluation performed by on authority and PBT assessment by another one), please tick the box indicating that this is a "Share substance".

On the website, only information agreed to be disclosed for this activity will be displayed. Nevertheless, it i advisable to provide more specific contact information (contact person, email) to ECHA in order to facilitat the communication between the organisation notifying an activity, other authorities and ECHA. Thi information will be kept internally in ECHA which will be in charge of facilitating communication an exchanges.

Identification of dossier submitter

information to be used on the ECHA website or formal communication when relevant.

Activity intended by: *

Please select country..

Please select country

Organisation name: *

Phone No:

Email:

Street

Street (cont):

City:

Region:

Postal Code:

Contact person - information for internal ECHA use

Title:

Please select title

First Name: *

Family Name: *

Email: *

Phone No:
(e.g. +358 111 222 333)

Shared substance:
More than one Authority is involved in this activity, the above organisation being the main authority responsible for this activity. Additional information can be provided in "Other remarks".

Information on the substance identification and composition

An activity can be performed for a substance with a well-defined composition (mono-constituent, multi constituent) or a substance of Unknown or Variable composition, Complex reaction products or Biologica materials (UVCB), or a group substance.

Show more...

Type of substance: *

Grouping approach

Substance (group) name: *

(For group substances you may provide name and identifiers of the representative (substance(s))

Per- and polyfluorinated substances (PFAS)

EC number: *

n.a.

CAS number (or other identification number): *

n.a.

Index number from CLP:

IUPAC name(s):

SMILES notation:

Other relevant substance information:

(Please use this field in case the Substance name/IUPAC name is longer that 255 characters)

For the purpose of this restriction intention PFAS are defined as fluorinated substances that contain at least one aliphatic carbon atom that is both, saturated and fully fluorinated, i.e. any chemical with at least one perfluorinated methyl group (-CF3) or at least one perfluorinated methylene group (-CF2-), including fluoropolymers and fluorinated side chain polymers.

General information

Does the substance contain other constituents/impurities/transformation products relevant for the activity?

Please add SID's as required

Add Constituent, Impurity, or Transformation product

(Potential) concern

Please specify here any of your (initial) concerns related to the substance that might be relevant for this or any of the follow-up processes. This helps to identify common areas of concern and improves coordination and consistency between the different processes.

In the below table, please tick the hazards relevant for your restriction proposal. Also tick use option, as this covers any (potential) use and is therefore relevant for restriction. Further details on the proposed restriction can be provided below.

Hazard/ Use Assessed
☑Carcinogenicity
☑Mutagenicity
☑Reproductive toxicity
☐Respiratory sensitisation
☐Skin sensitisation
☐Endocrine disruption for environment
☐Endocrine disruption for human health
☑Specific target organ toxicity - repeated
☐Other human health hazard
☑PBT/vPvB
☑Other environmental hazard
☑Use class covering all (potential) uses of the substance

Attachments

Attachments: Please add files relevant to your submission, such as dossiers, assessment documents, RCOMs, substance information or others. Please indicate in the filename if the file is confidential.

Add attachment | Browse

Maximum file size is 10 MB

If you would like to submit more than one document, please create a zip archive where you include all files and upload the zip file as attachment. Maximum file size is 100 MB.

Please note that if an access to documents application pursuant to Regulation (EC) No. 1049/2001 on public access to documents is received regarding this information, in order to define its position ECHA will have to first perform an assessment of the content of this information. Therefore, you are invited to indicate below any reason for which disclosure of this information should be denied. Please note that ECHA can only examine reasons appearing in Article 4 of the Regulation(EC) no 1049/2001 on public access to documents as valid.

I have the following reasons as set out in Article 4(1) or 4(2) of Regulation (EC) No 1049/2001 regarding public access to documents why the information submitted as confidential cannot be disclosed to persons requesting access to documents (please explain below in the commenting field those reasons; a reason could be that the protection of your commercial interests, including intellectual property, would be undermined).

The following information is relevant for this specific activity : Restriction

Please indicate if this is*

- ◉ An intention made according to Article 69(4) of REACH (this will be made public)
- ○ A pre-intention for the information of other MSCAs/ECHA/COM (this will not made public)

Scope and reason(s) for preparing a restriction dossier

the authority should indicate the scope, the target group(s) and the main reasons why it intends to prepare a restriction dossier, such as the main concern (human health, environment and/or human health via the environment), sources of exposures to be addressed (e.g. manufacture and/or all/specific use(s), including substances in articles).

Restriction on

(tick one or more options)

- ☑ Manufacturing
- ☑ Placing on the market
- ☑ Industrial use(s)
- ☑ Professional use(s)
- ☑ Consumer use(s)
- ☐ Other

Brief description of the intended restriction*

e.g. "Restricting the placing on the market of consumer articles containing substance x" or " Restricting the manufacturing of substance x and it compounds in consumer articles

Restriction on manufacture, placing on the market and use of PFAS

Main reason(s) for preparing a restriction dossier

All PFAS are, or ultimately transform into, persistent substances, leading to irreversible environmental exposure and accumulation. Due to their water solubility and mobility, contamination of surface, ground-, and drinking water and soil has occurred in the EU as well as globally and will continue. It has been proven very difficult and extremely costly to remove PFAS when released to the environment. In addition, some PFAS have been documented as toxic and/or bioaccumulative substances, both with respect to human health as well as the environment. Without taking action, their concentrations will continue to increase, and their toxic and polluting effects will be difficult to reverse.

Date of (pre)intention

15/06/2021

Expected date of submission*

15.07.2022

DD/MM/YYYY

Please note that for a Restriction these dates are informal. You will be asked at a later stage to give by e-mail the date of the final intention to be included in the RoI published on ECHA's website. This date will trigger the deadline of 12 months set out in the REACH regulation for submitting the Annex XV dossiers for restriction (Article 69(4)). Within this 12 months period you can define the date of submission, taking into account the dates as communicated by ECHA on its website.

Other remarks

Joint activity by Germany, the Netherlands, Denmark, Norway and Sweden.

Ich bin kein Roboter.

reCAPTCHA

Datenschutzerklärung - Nutzungsbedingungen

Note that fields are obligatory. If the information is not relevant, not available or unknown please record this accordingly.

The information submitted through this webform is used solely for the purpose of the relevant REACH process. The legal basis for the processing is found in Regulation (EC) No 1907/2006 on the Registration Evaluation Authorisation and Restriction of Chemicals (REACH).

The European Chemicals Agency will ensure on its part that your personal data is processed as required by Regulation (EC) No 45/2001 on the protection of personal data.

Any individual submitting personal data to ECHA is entitled to access and rectify that data. To exercise this right contact the data controller at annex-xv@echa.europa.eu. Furthermore, you also have the right to recourse to ECHA's Data Protection Officer (data-protection-officer@echa.europa.eu) or to the European Dat Protection Supervisor.

Submit to ECH