**Application** for exemption from requirements in Reach or CLP according to § 24 regulation (2008:245) concerning chemical products and biotechnological organisms

When relevant, safety data sheets for chemical products shall be submitted with the application.

1. **Information about the applicant (applicant organization and/or defense authority)**

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| --- | --- | --- |
| Applicant organization: | Business registration number: | Your Document ID: |
| Contact person: | Phone number: | E-mail: |
| The applicant’s role according to definitions in Reach and CLP:   * Manufacturer * Importer * Downstream user * Distributor * Other type of actor in the distribution chain | | |
| Brief description of the supply chain: | | |

1. **The exemption applies to**

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| --- | --- |
| * Exemption from *Regulation (EC) No 1907/2006 of the European parliament and of the Council concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals (Reach) and establishing a European Chemicals Agency* | Specify the article(s) of the regulations to which the application relates: |
| * Exemption from *Regulation (EC) No 1907/2006 of the European parliament and of the Council concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals (Reach) and establishing a European Chemicals Agency* | Relating to **article 33** |
| * Exemption from *Regulation (EC) No 1272/2008 of the European parliament and of the Council on classification, labeling and packaging of substances and mixtures (CLP)* | Specify the article(s) in the regulations that relate to the application |
| * Chemical product (go to section 3) | |
| * Complex object /article (go to section 4) | |

1. **Exemption regarding chemical product**

|  |  |  |
| --- | --- | --- |
| Trade name of chemical product (submit safety data sheet designed according to the Reach Regulation, Annex II): | | |
| Content of chemical product is:   * Known * Unknown | | |
| Enter chemical name, CAS number and classification of included substance(s) (alternatively submit an attachment with the data) | | |
| Chemical name: | CAS number: | Hazard classification: |
| The substance(s) are listed on the Candidate list: Yes ☐ No ☐  The substance(s) are listed on Annex XIV of the Reach Regulation: Yes ☐ No ☐ | | |

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| Classification of the product (specify hazard statements): |
| Area of use: |
| Estimated amount (per year): |
| Reason for applying for exemption due to the interests of the total defense (according to § 24(1)): |
| Describe how all possibilities to comply with the regulation have been exhausted: |
| Specify requests for the time period the exemption should apply and justification: |

1. **Exemption regarding complex object [[1]](#footnote-1)/article**

|  |  |  |
| --- | --- | --- |
| Name of the product (submit additional information if needed): | | |
| Enter the chemical name, CAS number and classification for the constituent substance/substances in the boxes below (alternatively attach an attachment with the data for all constituent chemical substances) | | |
| Chemical name: | CAS number: | Hazard classification: |
| The substance(s) are listed on the Candidate List: Yes ☐ No ☐  The substance(s) are listed on Annex XIV of the Reach regulation: Yes ☐ No ☐ | | |
| Enter the content and location of the substance(s) in the article[[2]](#footnote-2): | | |
| Enter the estimated amount of the substance(s) in the item and the number of articles/objects: | | |
| Reason for applying for exemption due to the interests of the total defense (according to § 24(1)): | | |
| Reason for application for exemption due to Swedish defense interests (according to § 24(2)): | | |
| Describe the purpose and justify why reporting and/or documentation to an external party according to current Reach and CLP cannot be fulfilled[[3]](#footnote-3): | | |
| Specify requests for the time period[[4]](#footnote-4) the exemption should apply and justification: | | |

1. **Assessment of risks**

Describe how the chemical product or article is to be used and the extent of the risks to health and the external environment, including waste management and storage, that are compatible with the use of the product or article. Also describe how these risks are managed and minimized.

1. **Applicant's signature**

|  |
| --- |
| Date: |
| Signature of the applicant: |
| Clarification of name and title/role: |

**Send the application either by post to the Inspectorate for Medicine and Environmental Health (FIHM), 107 85 STOCKHOLM, by e-mail to exp-fihm@mil.se or via VIDAR.**

**INFORMATION**

The exemption starts to apply when FIHM has made a decision on the matter, an approximate processing time is 3 months. If the form is incompletely filled out, FIHM will request a supplement to the case, which may delay processing.

**Information about the registration**

The information provided in this application will be processed by FIHM according to chapter 2 § 9 of the Defence Data Act in order for FIHM to be able to fulfill its supervisory responsibility. For more information about the Swedish Armed Forces’ processing of personal data and description of individuals’ rights, visit www.forsvarsmakten.se/personuppgifter.

FIHM have the capability of handling classified information. Prior contact is desired in these cases.

1. E.g., systems/complex objects/platforms. If this form is not sufficient because it is a very complex object, this entire description can be submitted in an appendix. [↑](#footnote-ref-1)
2. To ensure safe handling of product/article. The content applies to included articles that contain particularly dangerous substances if the substance is intentionally added and its weight content exceeds 0.1%. This is a legal requirement to report to Echa, regardless of whether the article is in a complex object or not. [↑](#footnote-ref-2)
3. For example, what activities the company has carried out to reach the conclusion that they are unable to meet the requirements. [↑](#footnote-ref-3)
4. If new substances are added to the Candidate list or Annex XIV during the time period, a new application regarding the new substances must be submitted. [↑](#footnote-ref-4)