Administrative information

This form shall be used by applicants submitting an application for authorisation of a substance intended to be used in materials and articles in contact with food. After completion it shall be submitted to the CBVV as part of the required information package.

The administrative information to be provided in this form relates to:

* Administrative data of the applicant submitting an application
* Subject of the request
* Existing authorisations

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| 1. **ADMINISTRATIVE DATA OF THE APPLICANT** |
| **1.1 Applicant[[1]](#footnote-1) (Company name):** |
| Telephone: |
| E-mail: |
| Address (street, number): |
| Post code: |
| City/Town: |
| Country: |
| **1.2 Name of contact person responsible for the application**: |
| Company: |
| Telephone: |
| E-mail: |
| Address (street, number): |
| Post code: |
| City/Town: |
| Country: |
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| 1. **SUBJECT OF THE REQUEST** |
| **2.1 Request for evaluation of a new substance:** |
| Answer 'yes' or 'no'.  If ‘no’ go to 2.2, if ‘yes’ give information requested in 2.1.1 to 2.1.4 as complete as possible. |
| * + 1. **Substance:** |
| * + 1. **CAS N°:** |
| * + 1. **Intended use/function:** |
| * + 1. **Inclusion in Dutch Commodities Act Regulation on packaging and consumer articles (Warenwetregeling verpakkingen en gebruiksartikelen):**   Please specify:  - Chapter and (sub)section of part A of the Regulation in which substance is to be listed  - Name and CAS No. under which it is to be listed  - Any restriction (e.g., SML) or specification to the listing |
| **2.2 Request for modification of an authorisation:** |
| Answer 'yes' or 'no'.  If ‘yes’ give information requested in 2.2.1 to 2.2.4 as complete as possible. |
| * + 1. **Substance:** |
| * + 1. **CAS N°:** |
| * + 1. **Inclusion in Dutch Commodities Act Regulation on packaging and consumer articles (Warenwetregeling verpakkingen en gebruiksartikelen):**   Please specify current entry |
| * + 1. **Reason for requesting the modification:** |
| **2.3 Request submitted also in other EU country/ies than NL:** |
| Answer 'yes' or 'no'.  If ‘yes’ specify country/ies where request has been submitted in addition to the Netherlands. |
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| 1. **AUTHORISATION OF SUBSTANCE** |
| **3.1 EU countries:** |
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| **3.1.1 In Member States:** |
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| **3.1.2 Notified as “new substance” in the context of Regulation (EC) No 1272/2008 on classification, labelling and packaging of substances and mixtures:** |
| If 'yes' give details and data transmitted: |
| **3.1.3 Other information:** |
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| **3.2 Non-EU countries** |
| **3.2.1 In USA:** |
| If 'yes' give relevant regulation(s) or other and give further details like restrictions and conditions: |
| **3.2.2 In Japan:** |
| If 'yes' give relevant regulation(s) or other and give further details like restrictions and conditions: |
| **3.2.3 In other countries:** |
| If 'yes' give relevant regulation(s) or other and give further details like restrictions and conditions: |
| * + 1. **Other information:** |
|  |
| **3.3 Other information:** |
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1. In case of more than one company submitting an application, their names and addresses should be provided. [↑](#footnote-ref-1)