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| --- | --- | --- | --- |
| **Annex 1 to the contract pursuant to DE-UZ 5** | | **Please use this printed form!** | |
| **Environmental label for “Sanitary Paper”** | |  | |
| Manufacturer (Applicant) |  | |
| Distributor (Label User) |  | |
| Brand/trade name: |  | |
| Product description[[1]](#footnote-1) (please tick the appropriate box) |  | |
| Crepe toilet paper (according to Paragraph 3.2) |  | |
| Crepe paper towels (according to Paragraph 3.3) |  | |
| Other sanitary paper products (according to Paragraph 3.4) |  | |
|  | (Please state the function e.g. toilet paper, facial tissues, etc.) | |
| Wet or dry strength agents added to the product and information on the quantity per 1 kg of dry pulp according to Paragraph 3.7 |  | |

Recovered paper grades according to Appendix 1 added to the finished product for the Basic Award Criteria DE-UZ 5:

|  |  |
| --- | --- |
| **Overview A** | |
| **Group / subgroups** | **Minimum % by mass** |
| 1 |  |
| 2 |  |
| 3 |  |
| 4 |  |
| 5 |  |
| Total in the finished product | 100 % |

Substances added during the manufacturing of the product according to   
Paragraphs 3.14 to 3.16

(please mark where applicable)

Colourant

Surface finishing agent

Auxiliary agents

Coating material

None

Bleaching agents and complexing agents added to the finished product according to   
Paragraph 3.17

|  |  |
| --- | --- |
| **Overview B** | |
| **Bleaching chemicals used** | **Complexing agents used** |
|  |  |

Biocidal substances (in preservatives and anti-slime agents) added to the product according to Paragraph 3.18

|  |  |  |  |
| --- | --- | --- | --- |
| Substance/trade name | Quantity per kilogram | IUPAC designation | CAS no. |
|  |  |  |  |

**Declarations by the applicant**

|  |  |  |
| --- | --- | --- |
| **Paragraph** | **We hereby declare that** | **Yes** |
| **3.1** | - the product is made from 100% recovered paper, |  |
| **3.2 – 3.4** | - the products according to Paragraphs 3.2, 3.3 and 3.4 comply with the requirements for the composition of the recovered paper and that the individual grades of recovered paper 2.14 and 4.07 are not added, |  |
| **3.5** | - the products do not exceed a maximum whiteness level of 80% (including the UV portion) according to DIN ISO 2470, |  |
| **3.6** | - only those process auxiliaries listed in Recommendation XXXVI from the BfR have been used and the maximum quantities and concentrations stated in this list have been observed, |  |
| **3.7** | - no wet or dry strength agents or other auxiliaries containing glyoxal are used in the manufacture of the sanitary paper products, |  |
| **3.8** | - optical brighteners have not been added, |  |
| **3.9** | - the concentration of chloropropanols in the water extract from the products manufactured using wet strength agents does not exceed the limits specified in Recommendation XXXVI from the BfR, |  |
| **3.11** | - the content of bisphenol A will be determined **once a year** for statistical purposes by an independent testing institution (certified according to ISO 17025) and submitted to RAL gGmbH, |  |
| **3.17** | - chlorine, halogenated bleaching agents and not readily biodegradable complexing agents such as e.g. ehylenediaminetetraacetic acid (EDTA) and diethylenetriaminepentaacetic acid (DTPA) have not been used in the processing of the recovered paper, |  |
| **3.18** | - only those substances that have been approved in accordance with the Biocidal Products Regulation (EU) 528/2012 (EU list of approved substances; formerly included in Annex I of the Biocidal Products Directive 98/09 EC) or that have been notified for the relevant biocidal product type and are still being tested as part of the EU review programme for existing active substances have been used as biocides, |  |
|  | - only those biocidal products that have been approved for their respective type of use have been used, |  |
|  | - products containing existing active substances that are still part of the EU review programme are still being used without approval until a decision has been reached, |  |
|  | - in addition, the product does not contain any substances that have been considered as candidates for substitution according to Article 10 of the EU Biocidal Products Regulation 528/2012, |  |
|  | - until the approval requirements for the respective biocidal products come into force, only those substances that are also listed in Recommendation XXXVI from the BfR are permitted and the following substances |  |
|  | * Sodium hexafluorosilicate CAS No. 16893-85-9 |  |
|  | * N-(α-(1-nitroethyl)benzyl) ethylenediamine CAS No. 14762-38-0 |  |
|  | * Tris-(hydroxymethyl)-nitromethane CAS No. 126-11-4, 5-chloro-2-methyl-4-isothiazolin-3-one, CAS No. 26172-55-4 and  2-methyl-4-isothiazolin-3-one CAS No.2682-20-4, |  |
|  | * Tetramethylthiuram disulfide CAS No. 127-36-8 |  |
|  | * Nano silver CAS No. 7440-22-4 |  |
|  | are not used, |  |
| **3.19** | - lotions, fragrances and bacterial suspensions are not used in the manufacture of the products, |  |
| **3.20** | - the products according to Paragraph 2 comply with the provisions of the German Food and Feed Code (Lebensmittel - Bedarfsgegenstände-und Futtermittelgesetzbuch (LFGB)). |  |

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| Please ***SELECT*** one answer (please only select **one** option) | | |
| **Paragraph** | **Furthermore, we declare that** | **Yes** |
| **3.18** | Colourants, surface finishing agents, auxiliaries and coating materials, which due to the use of -  2-methyl-4-isothiazolin-3-one or a mixture of  5-chloro-2-methyl-4-isothiazolin-3-one and -  2-Methyl-4-isothiazolin-3-one (3:1) have been labelled with the  **H-Phrase H317**, |  |
|  | have not been added |  |
|  | **or** |  |
|  | Colourants, surface finishing agents, auxiliaries and coating materials, which due to the use of -  2-methyl-4-isothiazolin-3-one or a mixture of  5-chloro-2-methyl-4-isothiazolin-3-one and -  2-Methyl-4-isothiazolin-3-one (3:1) have been labelled with the  **H-Phrase H317**, |  |
|  | have been added |  |
|  | These substances may be addedif it can be verified that the following quantities are not exceeded in total in the extracts from the finished products:   * Mixture of 5-chloro-2-methyl-4-isothiazolin-3-one, approx. 3 parts, and 2-Methyl-4-isothiazolin-3-one, approx. 1 part: 25 μg/dm² * 2-methyl-4-isothiazolin-3-one: 80 μg/dm²…”   The extracted quantities of the above-named biocides in the finished products have been verified in accordance with the “Guidelines for verifying the mass transfer from consumer goods made out of paper and board” (Leitfaden zur Überprüfung der Stoffübergänge von Bedarfsgegenständen aus Papier, Karton und Pappe) from the BfR.  The extracts were produced in accordance with the “Collection of methods for examining paper and board for food contact” (Methodensammlung zur Untersuchung von Papier, Karton und Pappe für den Lebensmittelkontakt) from the BfR and in accordance with DIN EN 645:1994-01, 647:1994-01 and 15519:2008-01. The quantity was determined using LCMS.  The test report from an independent testing institution accredited according to ISO 17025 or a testing institution recognised by the Federal Environmental Agency, e.g. Chair of Paper Technology and Mechanical Process Engineering at TU Darmstadt is enclosed. |  |

|  |  |  |  |
| --- | --- | --- | --- |
| **Location:** |  |  |  |
|  |  |  |
| **Date:** |  |  |

**Legally binding signature / company stamp**

1. This annex must be completed for each different product designation [↑](#footnote-ref-1)