



## Declaration of Compliance

|   |   |
|---|---|
| <b>Business Operator</b>                            | Vikan A/S<br>Rævevej 1<br>DK-7800 Skive<br>(+45) 96 14 26 00  |
| <b>Description</b>                                  | Hand Scoop, Metal Detectable, 1 Litre, Dark blue  |
| Item Number   | 567599  |
|   |    |
| Plastic Material                                    | Polypropylene with metal detectable agent   |
| <b>EU Compliance</b>                                |   |
| Regulation (EC) No 1935/2004                        | In accordance with EU Commission Regulation no. 1935/2004 article 3, 11(5), 15 and 17 the product is intended for food contact. The product is marked with the "glass & fork" symbol on the packaging or on the product itself through moulding.  |
|   |    |
| Regulation (EC) No 2023/2006                        | The product is produced according to EU Commission Regulation no. 2023/2006 of 22. December 2006 on good manufacturing practices for materials and articles intended to come into contact with food (GMP).  |
| Regulation (EU) No 10/2011                          | <p>Monomers and intentionally added additives used to manufacture this product are listed in Annex I of Commission Regulation (EU) No. 10/2011 of 14. January 2011 on plastic materials and articles intended to come into contact with foodstuffs. Subsequent amendments up to (EU) 2019/1338 are included.</p> <p>Monomers and/or additives with specific migration limit (SML) are used. The substances with a SML will not migrate in quantities that will exceed the SML, under the specified conditions of use. Upon request we will supply relevant information regarding these substances on a confidential basis.</p> <p>Vikan A/S does not use multi-layer materials or articles with a functional barrier.</p> |
| Regulations (EC) No 1333/2008 and (EC) No 1334/2008 | This material contains no "dual use" additives.   |
| <b>US FDA Compliance</b>                            |   |
|   | All raw materials in this product are in compliance with FDA (Food and Drug Administration in the USA) 21 CFR parts 170 to 199.   |
|   | The polymers and additives complies with FDA 21 CFR part 174, 175, 176, 177, 178, 181, 182, 184, or 186. Additives are cleared according to FDA 21 CFR Part 178 (Indirect food additives), are generally recognised as safe (GRAS), are prior-sanctioned food ingredients, or are cleared on basis of regulations for food additives of before 1958.  |
| <b>Danish Compliance</b>                            |   |
|   | The product complies with the Danish consolidation Act no. 1248 of 30/10/2018.  |

**Migration analysis plastics**

Samples of the product, or a similar product made from identical plastic material, have been tested for overall migration according to the test conditions specified in (EU) 10/2011, and the article comply with the overall migration limit of 10 mg/dm<sup>2</sup> or 60 mg/kg.

Test conditions for overall and specific migration were 5 min at 100 °C followed by 8 hours at 40 °C

Food simulants used for overall migration were 50 % ethanol (simulant D1), 3 % acetic acid (simulant B) and olive oil (simulant D2).

Compliance with specific migration limits, and other restrictions, has been documented through testing, calculation or simulation.

**Food contact types**

The product is suitable for contact with the following types of food under the intended and foreseeable conditions of use:

- Aqueous
- Acidic
- Alcoholic
- Fatty
- Dry

**Food contact usage time and temperature**

Any food contact conditions up to 100 °C

**Non-food contact usage temperature**

Minimum temperature: -20 °C  
Maximum temperature: 100 °C

**General**

Equipment should be cleaned, disinfected and sterilised, as appropriate to its intended use, before use.

It is also important to clean, disinfect and sterilise equipment as appropriate after use, using the appropriate decontamination chemicals, concentrations, times and temperatures.

Appropriate equipment decontamination will minimise the risk of microbial growth and cross contamination and will maximise the efficiency and durability of the equipment.

Recommended sterilisation temperature (Autoclave): 121 °C

We will make the relevant background documentation available to the competent authorities, at their request.

Vikan A/S is registered with the Danish Veterinary and Food Administration (DVFA), and our mandatory Own Control System is subject to inspection by the DVFA.

**Date**

05/09/2019



Made By

*Stine L. Bislev*

Stine Lønnerup Bislev  
Hygiene and Compliance Manager