

## **Declaration of Compliance**

Business Operator	Vikan A/S Rævevej 1 DK-7800 Skive (+45) 96 14 26 00	
Product name	Spare part Grip Band Module for 1011x, 1013x & HyGo, 5700x, Green	
Item Number	10032	
Plastic Material	Polypropylene, 98 % Thermoplastic elastomer (TPE), 98 %	
EU Compliance		
Regulation (EC) No 1935/2004	accordance with EU Commission Regulation no. 1935/2004 article 3, 11(5), 15 and 17 e product is intended for food contact. The product is marked with the "glass & fork" mbol on the packaging or on the product itself through moulding.	
	Ω,Ϊ	
AP(89)1	All pigments in the masterbatch comply with resolution AP 89(1)	
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	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) 2024/3190 are included.	
	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.	
	Vikan A/S does not use multi-layer materials or articles with a functional barrier.	
Regulations (EC) No 1333/2008 and (EC) No 1334/2008	This material contains intentionally added "dual use" additives for which restrictions or purity criteria are in place in accordance with Regulations (EC) 1333/2008 and (EC) 1334/2008. Upon request we will supply relevant information regarding these substances on a confidential basis.	

P (+45) 9614 2600 F (+45) 9614 2655 vikan@vikan.com www.vikan.com



US FDA Compliance	All raw materials in this product are in compliance with FDA (Food and Drug Administration in the USA) 21 CFR parts 170 to 199.			
	181, 182, 184, or 186. Additive food additives), are generally r	omplies with FDA 21 CFR part 1 es are cleared according to FDA recognised as safe (GRAS), are basis of regulations for food add	21 CFR Part 178 (Indirect prior-sanctioned food	
UK Compliance	The product complies with The Materials and Articles in Contact with Food (Amendment) (EU Exit) Regulations 2019 No. 704			
Danish Compliance	The product complies with the Danish consolidation Act no. 681 of 25/05/2020.			
Japanese Compliance	All substances (polymers, monomers and additives) used in Vikan products comply with Article 18(3) of the Japanese Food Sanitation Act and are listed in Tables 1 and 2 of Appendix 1 of the Positive List.			
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 for repeated use, and the article comply with the overall migration limit of 10 mg/dm <sup>2</sup> or 60 mg/kg.			
	Test conditions for overall migration were OM0 (30 min at 40 °C)			
	Food simulants used for overa acetic acid (simulant B).	Il migration were 50 % ethanol	(simulant D1) and 3 %	
	Compliance with specific migrathrough testing, calculation or	ation limits, and other restriction simulation.	is, has been documented	
Max ratio of food contact surface area to volume	e 2.1 dm²/100 ml			
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	o to 40 °C for 30 min.		
Non-food contact usage temperature	Minimum temperature: 0 °C Maximum temperature: 80 °C			
Vikan A/S CVR. 10290147	Rævevej 1 DK-7800 Skive	P (+45) 9614 2600 F (+45) 9614 2655	vikan@vikan.com www.vikan.com	



 General
 Equipment should be cleaned, disinfected and sterilised, as appropriate to it's 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.

Made By

Date

4/27/2025

A. Scholm

Marta Sztuka Materials and Compliance Specialist

P (+45) 9614 2600 F (+45) 9614 2655 vikan@vikan.com www.vikan.com