Avient Corporation
33587 Walker Road
Avon Lake, OH 44012

EU DECLARATION OF COMPLIANCE

Customer: Formerra
Product Code: EM1001609610
Product Name: DYNAFLEX™ G2711-1000-00
Certificate Date: 29 JAN 2026

It is hereby declared based on supplier information and/or own evaluation that the product DYNAFLEX™ G2711-1000-00 is a material that may be lawfully used in the manufacture of plastic articles that will contact food, subject to the restrictions provided below, in compliance with the harmonised legislation in place in the European Union (EU). Regulation (EU) 10/2011 on plastic materials and articles intended to come into contact with food is a specific measure within the meaning of Regulation (EC) No. 1935/2004. DYNAFLEX™ G2711-1000-00 meets the relevant requirements of European Union regulations 2023/2006 on materials and articles intended to come into contact with food.

DYNAFLEX™ G2711-1000-00 may contain polymers or additives in combination or singularly.

All additives that may be present in the product DYNAFLEX™ G2711-1000-00 are compliant with the relevant requirements of EU regulation 10/2011 as amended by EU regulations 321/2011, 1282/2011, 1183/2012, 202/2014, 865/2014, 2015/174, 2016/1416, 2017/752, 2018/79, 2018/213, 2018/831, 2019/37, 2019/988, 2019/1338, 2020/1245, 2023/1442, 2023/1627 and 2024/3190 on plastic materials and articles intended to come into contact with food (which repeals directive 2002/72/EC as amended).

Any polymers (excluding polymeric additives which are regulated as additives) that may be present in the product DYNAFLEX™ G2711-1000-00 are compliant with the relevant requirements of EU regulation 10/2011 on plastic materials and articles intended to come into contact with food, as amended by EU regulations 321/2011, 1282/2011, 1183/2012, 202/2014, 865/2014, 2015/174, 2016/1416, 2017/752, 2018/79, 2018/213, 2018/831, 2019/37, 2019/988, 2019/1338, 2020/1245, 2023/1442, 2023/1627 and 2024/3190 (which repeals directive 2002/72/EC as amended).

Applicable Restrictions (if any)

Components with SML	FCM223 ND This product contains at least one additional substance of Annex I and/or Annex II subject to SML*
Components with SML (T)	FCM294 5 mg/kg of food expressed as the sum of the substances and their oxidation products. Further substances to be considered see group restriction (14).
Components with QM	FCM223 1 mg/kg in final product.

This product contains at least one substance which is considered as dual-use additive (Article 11.3 & Annex IV of Regulation (EU) 10/2011).*

We are not aware of the intentional use of substances for which genotoxicity has not been ruled out. Based on the information from our raw material suppliers, PAAs not listed in Annex I are not expected to be present.

This product contains NIAS substance to be considered by downstream operator, as defined by regulation (EU) 1935/2004, Article 2*.

*For more information please contact your local representative. Disclosure may require a confidentiality agreement.

Please note that this product utilizes a white mineral oil (cleared for use in food and in food-contact articles) which does not fall under the mineral oil dosage limitation found within 21 CFR 177.2600. We also note that final extraction and migration limits may not be met by the finished part for fatty food contact applications.

General Conditions

The present certificate does not warrant against modifications of DYNAFLEX™ G2711-1000-00 resulting from its processing or from the addition of other products, nor against any inadequate use and/or storage of DYNAFLEX™ G2711-1000-00 or the materials and articles containing it.

Please note that article 12 of regulation 10/2011 specifies an overall migration limit of 10mg/dm2 or 60mg/kg.

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Extraction, migration, and NIAS testing, and organoleptic tests are the responsibility of the end-use manufacturer and are not part of our routine quality control. Unavoidable trace amounts of some elements listed in Annex II Table 1 of Commission Regulation (EU) 10/2011 may be present. As a producer of intermediate materials, Avient Corporation does not test for the migration of these elements in our products but these would be expected to contribute only marginally to any migration from a finished article.

With regards to the communication of intentional use of substances for which genotoxicity has not been ruled out, Annex I listed substances have not been considered as these will have been assessed as part of the EU 10/2011 petition process. Any applicable restrictions from Annex I are referenced above.

The use of this product manufactured by Avient Corporation or its subsidiaries and affiliates (Avient) in medical devices and drug packaging applications is not covered by the above or any other general regulation. It is the responsibility of the device or package manufacturer to establish safety with the FDA or other authorities through the submission of individual applications on the device or drug.

Avient does not control the conditions under which the products are used by the end-use manufacturer. Therefore, Avient does not warrant that the products meet all national regulatory requirements. It is the responsibility of the finished article manufacturer to ensure that all restrictions, including but not limited to food contact, including limits listed in Annex II Table 1 of Commission Regulation (EU) 10/2011, and organoleptic requirements, are met by the finished article and that such article is fully compliant with all the relevant national regulations.

The information provided pursuant to current regulations is valid as of the date of this letter and from one-year following its issuance or until changes of the cited regulations become effective whichever comes earlier. Avient makes no representations or warranties as to the accuracy of the information contained or referenced as of any other date. This certificate applies to the product manufactured in North America.

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