

Declaration concerning food contact

FDA (Food & Drug Administration)

Product

Exolon® GP colourless

We hereby declare that our above-mentioned semi-finished product, due to the resin manufacturer's data, complies with the requirements of the FDA regulation 21 CFR, Part 177, Section 1580 „Polycarbonate resins“. Any additives that may be present comply with appropriate, specific FDA regulations.

Restrictions

Conditions of use: The material is suitable for contact with all food types under Conditions of Use B to H as per FDA.

Additional: Finished food contact articles made from this material are subject to an extractive restriction in accordance with FDA regulation 21 CFR 177.1580.

Compliance with these requirements was not checked on the semi-finished product, as these generally relate to the final article. The exact provisions of the above-mentioned FDA regulation can be found at:
<https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfcfr/cfrsearch.cfm>

The quality assurance system of our raw material supplier is certified as per DIN EN ISO 9001:2015 and serves as an important basis of the constant composition and quality of Exolon® GP colourless.

Note:

It is the customer's own responsibility to test the suitability of plastic items manufactured from or with our products for the planned application in the foodstuff industry. That includes for example:

- testing, if the physical characteristics of the plastic are suitable for the planned application,
- testing, if plastic parts manufactured by the customer fulfil the prescribed migration or extraction values,
- testing for possible influence of the plastic on the composition and/or organoleptic characteristics of foodstuffs.

This information is based on the information provided by our suppliers. We are not liable for the completeness and correctness of information contained herein. Food-contact suitability depends on the grade and must be confirmed for the specific type used. Existing laws and regulations must be respected by the receivers/users of our product in their own responsibility.

All information contained in this document is provided in good faith and is based on sources believed to be reliable at the time of publication of this document. In the event of changes, for example due to legislation, manufacturing-related changes, or new scientific findings, this declaration will be reassessed and becomes invalid. This declaration expires automatically 12 months after the date of issue. It is the sole responsibility of our customer to ensure that the laws and regulations necessary for their intended use are complied with. Please request an up-to-date confirmation from Liedtke Kunststofftechnik if required.

The Food and Drug Administration (FDA) is an agency of the United States Department of Health and Human Services, one of the United States federal executive departments.

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