Liedtke Kunststofftechnik
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Food Contact Declaration

FDA (Food & Drug Administration)

Product: Perlon® (PA 6) natural

We hereby declare that our semi-finished products made of the above product, with regard to their chemical composition and according to the information provided by the raw material manufacturers, comply with the requirements of FDA regulation 21 CFR, Part 177, Section 1500 "Nylon resins".

Restrictions

- **Scope:** Sheets from 6 mm to 100 mm / round rods Ø 20 mm to Ø 200 mm.
- **Other:** The finished food-contact article is subject to the extraction restriction under FDA regulation 21 CFR 177.1500 (a)(6) (Nylon Resins).

Compliance with these requirements has not been tested on the semi-finished product, as they generally relate to the finished article. The exact provisions of the above FDA regulations can be found at: https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfcfr/cfrsearch.cfm

Our raw material supplier's quality management system is certified to DIN EN ISO 9001:2015 and forms an important basis for the consistent composition and quality of Perlon® (PA 6) natural.

Perlon® is a registered trade name for polyamide 6; the above declaration refers to the chemical composition of the base polymer polyamide 6.

General note:

It remains the customer's responsibility to verify the suitability of the plastic articles made from or with our products for the intended food-contact application, i.e. for example:

- checking whether the physical properties of the plastic are suitable for the intended applications,
- checking compliance with migration or extraction limits on the finished plastic parts,
- checking the possible influence of the plastic on the composition and/or organoleptic properties of foodstuffs.

The Food and Drug Administration (FDA) is the food and drug regulatory authority of the United States, under the Department of Health and Human Services.

This information was generated automatically and therefore bears no signature.

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