

Food Contact Legal Declaration

FDA (Food & Drug Administration)

Product: PF CP 206 (Hp 2062.8)

We hereby declare that the resin component (phenolic resin) of the product listed below, according to the raw material manufacturers' information, complies with the requirements of FDA regulation 21 CFR, Part 177, Section 2410 "Phenolic resins in molded articles".

Restrictions

- **No suitability for food contact:** PF CP 206 (Hp 2062.8) is a technical electrical-insulation phenolic paper laminate (paper substrate, phenolic-resin-bonded) and as a finished laminate is NOT declared suitable for direct food contact.
- **Other:** No testing for extraction limits under FDA regulations has been carried out on the finished laminate, as the product is intended for technical insulating and punching applications (electrical, switchgear, mechanical engineering).

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 to DIN EN ISO 9001:2015 and forms an important basis for the consistent composition and quality of PF CP 206 (Hp 2062.8).

General note:

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

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

Our statements are based on documents provided by our suppliers. No liability is accepted for the completeness or accuracy of the information contained herein. Applicable laws and regulations must be observed by the recipient/user of our products at their own responsibility.

All information contained in this document is provided to the best of our knowledge and is based on sources considered 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 become invalid. This declaration automatically expires 12 months after the date of issue. It is the sole responsibility of our customer to ensure compliance with the laws and regulations necessary for their intended use. Please request an updated confirmation from Liedtke Kunststofftechnik if required.

The Food and Drug Administration (FDA) is the food safety and drug approval authority of the United States and is subordinate to the Department of Health and Human Services.

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