Prohlášení týkající se styku s potravinami

FDA (Úřad pro kontrolu potravin a léčiv)

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Produkt

Sustamid® 66 přírodní

We hereby declare that our above-mentioned semi-finished product, due to the resin manufacturers data, comply with the requirements of the FDA-regulation **21 CFR, část 177, oddíl 1500 „Nylonové pryskyřice“ týkající se jejich chemického složení, jakož i dalších specifických předpisů FDA pro zabudované přídatné látky.**

Omezení

Další: Na hotový výrobek určený pro styk s potravinami se vztahuje omezení extrakce podle nařízení FDA 21 CFR 177.1500.

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 aboved-mentioned FDA-regulation can be found at:
https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfcfr/cfrsearch.cfm

The quality assurance system of Röchling Industrial Lahnstein SE & Co. KG is certified as per DIN EN ISO 9001:2015 and serves as an important basis of the constant composition and quality of Sustamid® 66 přírodní.

Note:

It is customers 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 fulfill the prescribed migration or extraction values,
- testing for possible influence of the plastic on the composition and/or organoleptic characteristics of foodstuffs.

This information above is based on the information provided by our suppliers. We are not liable for completeness and correctness of information contained herein. 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, new statements will be published on our websitehttps://www.roechling.com/industrial/materials/. Previous declarations will become invalid as a result. This declaration expired 12 months after the date of issue (Print). It is the sole responsibility of our customer to ensure that the laws and regulations necessary for their intended use are complied with. Therefore, if necessary, please request a new confirmation or download it from our website https://www.roechling.com/industrial/materials/.

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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