PHACOTEC
PRODUKT-SERVICE GMBH
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DEUTSCHLAND

Declaration of Conformity (PPWR)

Series	Standard Pharma-Tech
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This document was created digitally based on information provided by the responsible company (1) and is valid without a signature.

This document is provided for information purposes only. Only the current original manufacturer documentation and declarations of compliance issued by the respective manufacturer shall be considered binding. In the event of discrepancies, the manufacturer documentation shall prevail.

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Supplementary Notes and Explanations regarding Regulation (EU) 2025/40

(1) General
The present PPWR compliance assessment is based on manufacturer information, raw material specifications, supplier declarations, test reports and other technical documentation available at the time of document preparation.
Unless expressly stated otherwise, the information has not been independently verified by the distributor.

(2) Technical Documentation
The technical documentation required under Article 39 of Regulation (EU) 2025/40 is maintained by the respective producer or manufacturer.
Access to complete technical documentation may be restricted due to trade secrets and confidential business information.

(3) Product Families and Material Groups
Manufacturer declarations, technical assessments and compliance evaluations are generally based on material families, product groups, formulations, material/colour combinations or worst-case evaluations.
Accordingly, such documentation generally applies to all covered article variants, provided that material composition, manufacturing process and intended use remain identical.

(4) Recyclability (Article 6)
The recyclability assessment is based on the legal requirements, manufacturer information and recognised evaluation methods available at the time of document preparation.
Should the European Commission publish binding harmonised assessment methodologies or delegated acts in the future, reassessment may become necessary.

(5) Recycled Content (Article 7)
Statements regarding recycled content are based on manufacturer information and available raw material documentation.
Unless expressly stated otherwise, no independent material analyses have been performed to determine recycled content.

(6) PFAS
Statements regarding PFAS are based on manufacturer and supplier declarations as well as available raw material specifications.
Analytical testing for individual PFAS substances, Total Fluorine (TF), Total Organic Fluorine (TOF) or comparable screening methods is only available where such testing has been expressly performed by the respective manufacturer or an accredited laboratory.

(7) Bisphenols
Statements regarding Bisphenol A (BPA), other bisphenols and bisphenol derivatives are based on manufacturer and supplier declarations as well as available raw material information.
Unless otherwise documented, no independent analytical testing for BPA, BPF, BPS or other bisphenol compounds has been performed.

(8) REACH and SVHC
The assessment regarding Regulation (EC) No. 1907/2006 (REACH) and the Candidate List of Substances of Very High Concern (SVHC) is based on the manufacturer information available at the time of document preparation.
Updates to the Candidate List may require an update of the underlying manufacturer declarations.

(9) Heavy Metals
Statements regarding lead, cadmium, mercury and hexavalent chromium are generally based on raw material approvals, supplier declarations, material specifications or manufacturer test reports.
Product-specific laboratory analyses are generally not available for each individual article number.

(10) Substance Testing and Product Families
Analytical testing regarding PFAS, heavy metals, migration limits or other substance restrictions is generally performed by manufacturers on material, formulation, colour or product-family level.
Separate testing for each individual article number, colour variant or production batch is generally not performed unless expressly agreed otherwise.

(11) Labelling (Article 12)
Where transitional provisions, guidance documents, implementing acts or technical specifications issued by the European Commission are still pending for specific packaging categories, implementation will take place once such requirements enter into force.

(12) Basis of Documentation
The present assessment has been prepared on the basis of manufacturer information available at the time of issue.
Updated manufacturer documentation or changes in legal requirements may require revision of the assessment.

(13) Disclaimer
This annex is provided for information purposes only.
Only the current declarations, technical documentation and compliance evidence issued by the respective producer or manufacturer shall be considered binding.
In the event of discrepancies, the original manufacturer documentation shall prevail.