

Certificate of conformity BSE / TSE

Material types:

Ceradur	PA 6 G + Oil	PTFE	St 6000 [®] ELS FDA
CeramX [®]	PA 6 G + Oil blue	PVC	St 6000 [®] GB
Flex 11	PA 12 G	PVDF	St 6000 [®] MDP
Flex 77	PEEK	St 1000 [®]	St 6000 [®] XDP
Flex 88	PETP	St 4000 [®] ATEX	St 7000 [®] AMB
Flex Line [®]	PETP-SP	St 500 [®]	St 7000 [®] EHT
Flex Wear	POM	St 6000 [®] AST	St 9100 Oil
CeraFlex	PP	St 6000 [®] AST	
PA 6 G	PS 1000 [®]	FDA	
		St 6000 [®] ELS	

Based on the information provided by our preliminary supplier we confirm to you that the products made of the materials listed are perfectly sound according to the directive concerning protective meat hygiene measures to combat bovine spongiform encephalopathy (= BSE directive) and transmissible spongiform encephalopathy (= TSE directive).

During production, storage and shipping, the material does not come into contact with contaminated or non-contaminated meat as defined by the directive. A routine check on the presence of all these materials is not carried out at the exit inspection. We also confirm that according to our preliminary supplier, only new material is used.

This declaration does not relieve the issuer from the obligation to check the goods for their own purposes.

General notes:

The declaration reflects the latest technological and scientific standards as of the date specified and is not equivalent to a guarantee. No responsibility is accepted for the completeness and accuracy of the information contained. The recipient / user of our products is responsible for adhering to current laws and regulations. Our declaration is based on documents we have received from our suppliers and on calculations and migration tests carried out by third parties, where relevant.

As soon as the law / directive / guideline, raw material, formulas, handling processes or similar are changed, the declaration has to be updated (no later than every three years). Previous versions automatically become invalid as a result. We recommend requesting the declaration at regular intervals.

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Since Wefapress Beck + Co. GmbH does not have any information regarding the conditions of use (such as contact medium, contact time and temperature as well as hygienic conditions), the distributing party (plant manufacturer or operator) must carry out the final and conclusive test with reference to the information on migration limits provided.

