



Customer Assurance Statement

TRANSMITTABLE / BOVINE SPONGIFORM ENCEPHALOPATHY (TSE/BSE)

This statement refers to all products supplied by any of the Ingredion EMEA Companies.

The products which the Ingredion EMEA Company supplies to you are not affected by the "Note for Guidance on Minimising the Risk of TRANSMITTING Animal Spongiform Encephalopathy (TSE) via Medicinal Products" referred to in the Annex to Directive 75/318/EEC as amended by Directive 1999/82/EEC¹ concerning TSE or the General Framework Regulation (EC) 999/2001² and its amendments, laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies, which is known as the "TSE Regulation".

Please note, the information given in this statement is in relation to products supplied by any of the Ingredion EMEA Companies and is based upon their interpretation of relevant legislation. Although it is offered in good faith, the advice is not legal advice to you. It is therefore necessary that you satisfy yourself of the use and any labelling obligations, in accordance with relevant legislation, for your products as sold to the ultimate consumer. Each of the Ingredion EMEA Companies cannot accept any liability in this regard. The 'Ingredion EMEA Companies' are each of Ingredion UK Limited, Ingredion Germany GmbH, Ingredion Middle East Branch, Ingredion South Africa Pty Limited and Ingredion Holding LLC- Kenya Branch Office. Issued on behalf of each of the Ingredion EMEA Companies by Ingredion UK Limited.

TSE/BSE Statement

July 2017

¹ Commission Directive 1999/82/EC of 8/9/99, amending the Annex to Council Directive 75/318/EEC on the approximation of the laws of the Member States relating to analytical, pharmacotoxicological and clinical standards and protocols in respect of the testing of medicinal products.

² Regulation (EC) No 999/2001 of 22 May 2001 laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies