



## Customer Assurance Statement

# TRANSMITTABLE / BOVINE SPONGIFORM ENCEPHALOPATHY (TSE/BSE)

This statement refers to all products, excluding Food Systems Solutions<sup>1</sup>, supplied by any of the Ingredion EMEA Companies.

The products which the Ingredion EMEA Companies supply to you are not affected by the Note for Guidance on Minimising the Risk of Transmitting Animal Spongiform Encephalopathy Agents via Medicinal Products<sup>2</sup> (as referred to in Directive 2001/83/EC on the Community code relating to medicinal products for human use)<sup>3</sup>.

Furthermore, these products are not affected by the General Framework Regulation (EC) No 999/2001<sup>4</sup> 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

January 2021

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<sup>1</sup> Statement available on request

<sup>2</sup> Note for guidance on minimising the risk of transmitting animal spongiform encephalopathy agents via human and veterinary medicinal products (EMA/410/01 rev.3) (2011/C 73/01)

<sup>3</sup> Directive 2001/83/EC of the European Parliament and of the Council of 6 November 2001 on the Community code relating to medicinal products for human use

<sup>4</sup> Regulation (EC) No 999/2001 of the European Parliament and of the Council of 22 May 2001 laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies, as amended