

Safety assessment Minca Supplement

According to Regulation (EC) No. 1907/2006
Issue date: 2014-10-01, Replaces version: XXXX-XX-XX



Article number

TN 1334

Trade name

Minca Supplement

Intended use

Encourage the development of the fimbrial antigen (F5 (K99)) in pathogenic *Escherichia coli* together with Minca Agar, modified, REF TN 1722 and TN 1040. Relevant to veterinary medicine (young cattles).

Name of supplier

sifin diagnostics gmbh
Berliner Allee 317-321
13088 Berlin, Germany

Contact for technical information

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E-Mail (competent person)

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Emergency telephone number

Phone: +49 30 927 030-0, Safety Officer for Medical Devices
Mo. - Th. 7.30 a.m. - 4.15 p.m., Fr. 7.30 a.m. - 3.00 p.m.

Classification of the mixture

Classification according to Regulation (EC) 1272/2008 (CLP): This mixture is classified as not hazardous according to Regulation (EC) 1272/2008 (CLP).

Classification according to Directive 1999/45/EC (DPD): This mixture is classified as not hazardous according to Directive 1999/45/EC.

Label elements

Labelling according to Regulation (EC) 1272/2008 [CLP]: Not applicable.

Labelling according to Directive 1999/45/EC (DPD): Not applicable.

Other hazards

None known. Minimum standards for protection measures (TRGS 500) as common in the chemical industry should be followed.

According to Article 31 of the REACH Regulation, drawing up a safety data sheet for this product is unnecessary, as the product is neither classified as a hazardous material nor does it contain hazardous ingredients or materials with Community (EC) workplace exposure limits or substances of very high concern (SVHC) above their particular legal nominal limits. Under REACH Regulation (EC) no. 1907/2006 (REACH), no safety data sheet is therefore required and also not available in this case.