7002 S. 109th STREET, LA VISTA, NE 68128

IMPORTANT SDS INFORMATION FOR ARM-D® Kit, TEM/SHV/GES

Dear Customer:

The above product is manufactured by Streck and does not meet the criteria for classification as hazardous in accordance with Regulation (EC) No 1272/2008 of the European Parliament and of the Council of 16 December 2008 on classification, labelling, and packaging of substances, Directive 98/79/EC on In Vitro Diagnostic Medical Devices and mixtures, REACH Regulation (EC) No 1907/2006 and Commission Regulation (EU) 2021/979 amendment; or where a substance is persistent, bioaccumulative and toxic or very persistent and very bioaccumulative in accordance with the criteria set out in Annex XIII.

Product complies with all applicable regulations of the U.S. Department of Transportation (DOT) and/or International Air Transportation Association (IATA), and does not meet criteria for restricted transport.

Streck's products are classified as *in vitro* diagnostics and regulated by the FDA. Accordingly, all applicable precautions are stated on the product label or package insert that accompanies each product.

Streck suggests you place a copy of this letter in any place you would ordinarily keep an SDS. This step will document your laboratory's compliance with OSHA requirements.

Streck Safety Administrator