



Test report

Skin Irritation Test

Test report Number: ISO 201708-02242C_SI_engl

commissioned by:

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Sep 20th 17

Test material:

EVA-plastic „EVA SPEZIAL SANDWICH TRIPLEX®“

Test material received: Aug 16th 17

Test performed: Sep 15th 17

<u>Result</u> The EVA-plastic „EVA SPEZIAL SANDWICH TRIPLEX®“ did not cause a skin irritating effect.

Description of the test procedure:

Normative References: ISO 10993-10 (2009), ISO 10993-1 (2009), OECD TG 439

Based on the normative references ISO-10993-1 (2009), Chapter 4.6 and ISO 10993-10 (2009) an *in vitro* test was performed to evaluate the potential skin irritating effect of the test material. An *in vitro* reconstituted, human epidermal 3D-skin model, type epiCS, Lot. No. 100-AG1636-1 was used. The test was performed according to ECVAM's "Performance standards for applying human skin models to *in vitro* skin irritation testing".

Prior to performing the test procedure the skin model cell culture inserts were incubated for 24 h at 37°C and 5 % pCO₂ in a cell culture incubator in fresh Maintenance Medium. After this preincubation time 50 µl of PBS was pipetted on the surface of each skin model. Then the material samples to be tested were applied on the skin surface using a pair of sterile tweezers (insert surface diameter approximately 8 mm)

50 µl Triton X 100 were used as a (skin irritating) positive control, 50 µl PBS were used as a (not skin irritating) negative control. All experiments were performed in duplicate. After 20 min incubation all inserts were rinsed thoroughly with sterile PBS, blotted on a paper towel to remove excess PBS and cultivated in Maintenance Medium for 42 h in a cell culture incubator at 37°C and 5 % pCO₂ .

Measurement of LDH-release:

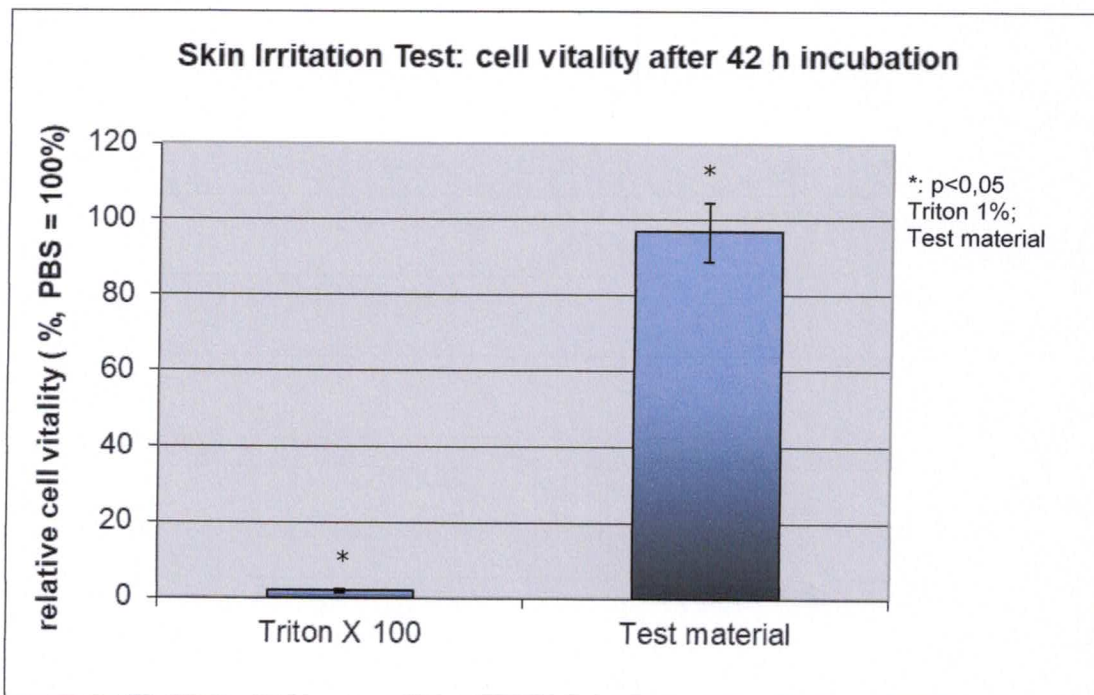
After the 42 h incubation period 2 x 100 µl Maintenance Medium Samples were taken from every skin model and the LDH-release was measured using a „Cytotoxicity Detection Kit (LDH)“ from Roche Diagnostics.

Measurement of cell vitality (MTT-test):

The inserts were rinsed once in Assay Medium using a 24-well cell culture plate and transferred in a second 24-well cell culture plate with 300 µl Cellsystems Assay Medium containing 1mg/ml MTT (Sigma M5655). All inserts were incubated for additional 3 h in a cell culture incubator at 37°C and 5 % pCO₂. Afterwards all inserts were blotted on a paper towel and the MTT-dye was extracted from the skin samples using 2 ml Isopropanol per insert. The extinction of each Isopropanol extract was measured in a photometer at 570 nm. With this data the relative vitality of the cells in the skin samples compared to the negative control (PBS) was calculated.

Results:

Measurement of cell vitality (MTT-test)

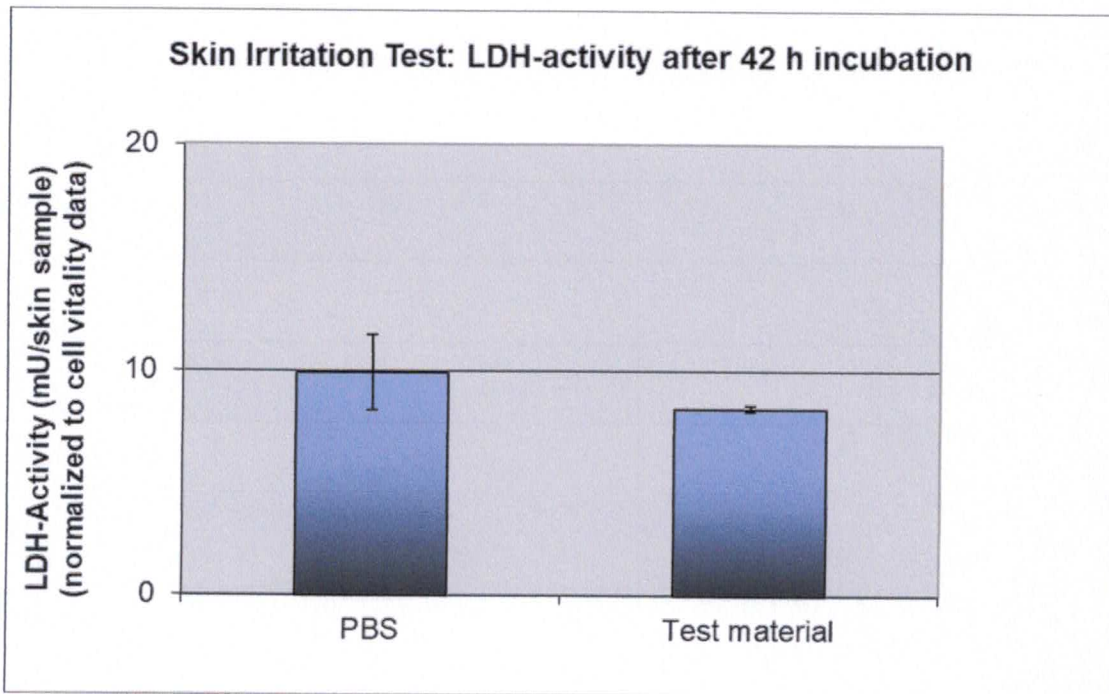


Result data (rel. cell vitality)	n=2, (PBS data not shown in graphic)		
	Triton X 100	PBS	Test material
Mean	1,93 %	100 %	96,74 %
Standarddev.	0,42 %	8,31 %	7,87 %

In the presence of Triton X 100 on the skin culture inserts 1,9 % of the cell vitality compared to the negative control was reached. This value is within the valid range of 15 % cell vitality or less compared to the negative control.

A material is considered as skin irritating, if it reduces the cell vitality of the skin samples to less than 50 % compared to negative control skin samples. This is not the case in this experiment. The material did not show a skin irritating effect.

LDH-Release



LDH-Release (mU/skin sample)	Triton X 100 data not shown in graphic, n=2		
	Triton X 100	PBS	Test material
Mean	4252,86	9,94	8,36
Standarddev.	42,27	9,07	0,16

Skin samples charged with the test material did not show a significant difference in LDH-release compared to the PBS negative controls.

Result **The EVA-plastic „EVA SPEZIAL SANDWICH TRIPLEX®“
did not cause a skin irritating effect.**

Explanatory notes:

none

Test performed by: *Dietrich Scheddin*

authorized by: *Dietrich Scheddin*
(Dr. D. Scheddin / CEO CYTOX)

It is not allowed to publish only parts of this test report without written approval of CYTOX.