Test report number: ISO 201708-02242A_SI_engl

Test report

Skin Irritation Test

Test report Number: ISO 201708-02242A_SI_engl

commissioned by:

W.R. Lang GmbH Hafenstraße 83 D-56564 Neuwied, GERMANY



CYTOX biological testing of medical devices Gottlieb-Keim-Straße 60 95448 Bayreuth, Germany

tel. +49-921-1511-254 fax +49-921-1511-255 mobil +49-179-5102577

info@cytox.de www.cytox.de

Sep 20th 17

Test material:

EVA-plastic "LaNe® EVA SKIN"

Test material received: Aug 16th 17

Test performed: Sep 15th 17

Result The EVA-plastic "LaNe® EVA SKIN"

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 % pCO $_2$ in a cell culture incubator in fresh Maintenance Medium. After this preincubation time 50 μ l 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)

CYTOX biological testing of medical devices

 $50~\mu l$ Triton X 100 were used as a (skin irritating) positive control, $50~\mu l$ 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 $37^{\circ}C$ and 5~% pCO $_2$.

Measurement of LDH-release:

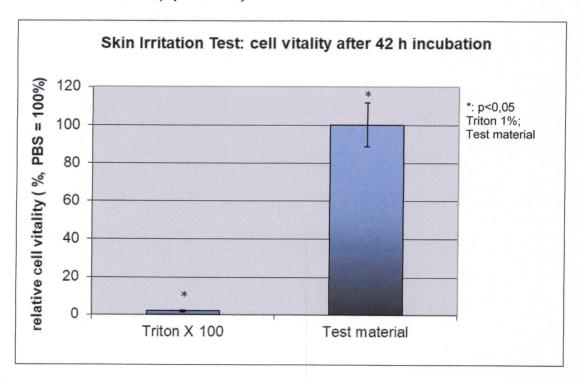
After the 42 h incubation period 2 x $100~\mu$ l Maintenance Medium Samples were taken from every skin model and the LDH-release was measured using a "Cytotoxicity Detection Kit (LDH)" from Roche Diagnostics.

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 μ l 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)



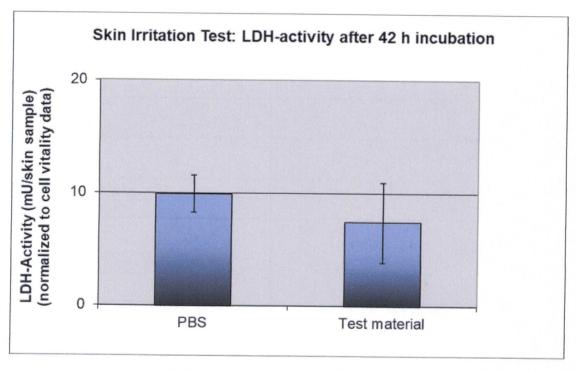
Result data	n=2, (PBS data not shown in graphic)			
(rel. cell vitality)	Triton X 100	PBS	Test material	
Mean	1,93 %	100 %	100,22 %	
Standarddev.	0,42 %	8,31 %	11,43 %	

CYTOX biological testing of medical devices

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	7,42
Standarddev.	42,27	9,07	3,60

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

Test report number: ISO 201708-02242A_SI_engl

Result The EVA-plastic "LaNe® EVA SKIN" did not cause a skin irritating effect.

Explanatory notes:

none

Test performed by: Dita Schoolston

authorized by: Dat S. Ladden (Dr. D. Scheddin / CEO CYTOX)

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