Department : Cell Biology

STUDY NO: RR222098/CB/CP/09-22



DRAFT REPORT

Copy No. 1/2

Study Title

Evaluation of *In vitro* Cytoprotective effect of test substance against UV induced cell damage in Human Keratinocytes (HaCaT) cell line

Study Director

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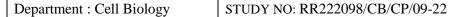
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COMPLIANCE STATEMENT

The Study Director hereby declares that the work was performed under his supervision and in accordance with the mutually agreed study plan and the in house procedures. It is assured that the reported results represent the raw data obtained during the experimental work. No circumstances have been left unreported which may have affected the quality or integrity of the data or which might have a potential bearing on the validity and reproducibility of this study. The Study Director accepts overall responsibility for the technical conduct of the study as well as the interpretation, documentation and reporting of the results.

Date: 05/11/2022 Study Director
Dr. Ashok G





CERTIFICATE OF AFFIRMATION AND CONFIDENTIALITY

The Management hereby attests to the originality, accuracy and authenticity of the study to the best of their knowledge. This report contains confidential and proprietary information of **M/s.Tocyen Beauty Cream, Mumbai, India.,** which will not be disclosed to anyone without the expressed or written approval of authorized personnel.

Date: 05/11/2022

Management Dr. Ashok G C.E.O

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DECLARATION

The Study No. RR222098/CB/CP/09-22, entitled "Evaluation of *In vitro* Cytoprotective effect of test substance against UV induced cell damage in Human Keratinocytes (HaCaT) cell line" has been inspected regularly according to the Standard Operating Procedure of the test facility's Quality Assurance Unit. The report was audited against approved study plan and pertinent raw data and accurately reflects the raw data.

Date: 05/11/2022 QA Head Mr. Gopi M

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ABBREVIATION USED

MCR : Microbiology % : Percentage

CB : Cell Biology gm : Gram

MB : Molecular Biology hr : Hour

BC : Biochemistry mg : Milligram

DTL : Drug Testing Laboratory mL : Milliliter

PC : Preclinical nm : Nano meter

CL : Clinical μL : Microliter

PBS : Phosphate buffer saline μg : Microgram

°C : Degree Centigrade

EDTA : Ethylenediaminetetraacetic acid

RT-PCR : Reverse transcription-polymerase chain reaction

dNTP : Deoxynucleotide Triphosphate

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LIST OF TABLE

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1.	In vitro Cytoprotective effect of test product in Human Keratinocytes (HaCaT) cell line against UV induced cell damage.	12



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1. STUDY DETAILS

1.1. Study title : Determination of *in vitro* cytoprotective of

test product against UV induced cell damage

in Human Keratinocytes (HaCaT) cells.

1.2. Study number : RR222098/CB/CP/09-22

1.3. Test Product : Tocyen Beauty Cream

1.4.Sponsor : M/s. Tocyen Beauty Cream,

Mumbai, India.,

1.4. Test Facility : Radiant Research Services Pvt. Ltd.

99/A, 8th Main, 3rd Phase,

Peenya industrial area,

Bangalore-560058

1.5. Test schedule

Study Initiation Date : 14/09/2022

Experimental Start Date : 16/09/2022

Experimental Completion Date : 03/11/2022

Study Completion Date : 05/11/2022

1.6. Study Responsibilities

Study Director : Dr. Ashok G

Study coordinator : Mr. Vatsa Kapadia

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2. OBJECTIVE

The purpose of this study is to evaluate the *in vitro* cytoprotective activity of test substance against UV induced damage in Human Keratinocyte (HaCaT) cell line.

3. SUMMARY

The Tocyen Beauty Cream was evaluated for its *in vitro* cytoprotective activity in Human Keratinocyte cell line (HaCaT) cell line, the test product was first evaluated for its cytotoxicity with different concentrations from 1000μg/mL – 7.8μg/mL. The test product exhibited a CTC₅₀ value above 1000μg/mL on the Human Keratinocyte (HaCaT) cell line. Hence non-toxic concentrations were taken for further study.

In the *in vitro* cytoprotective study, the test product at tested concentrations (1000µg/mL and 500µg/mL) showed cytoprotective activity against UV induced damage.

4. GUIDELINES/REFERENCE

1) Francis D and Rita L. "Rapid colorometric assay for cell growth and survival modifications to the tetrazolium dye procedure giving improved sensitivity and reliability". Journal of Immunological Methods, 1986; 89: 271-277.

5. AMENDMENT AND DEVIATION PROCEDURE

No deviation has been adapted during the conduct of the experiment.

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6. MATERIALS

6.1. Test product information

Sample code	Sample name	Name used in the report	Batch number	Physical appearance	Storage condition
RR222098	Tocyen Beauty Cream	RR222098 / Tocyen Beauty Cream	CG6122	Semi-solid	RT

6.2. Reference Material/Chemicals

Chemicals	Batch / Lot No.	Manufacturer	Expiry Date
MTT	0000454015	HiMedia, India	Oct-2024
DMEM-HG	2365585	Gibco, USA	Feb-2024
Fetal Bovine serum	2422662	Gibco, Brazil	Sep-2026
DPBS	0000474192	HiMedia, India	Mar-2024
Trypsin - EDTA	0000472777	HiMedia, India	Mar-2023
Antibiotics	0000493509	HiMedia, India	Aug-2023
DMSO	J19A/0416/0305/13	SDFCL, India	Sep-2024

6.3. Equipments

S. No.	Name of the Instrument	Make	Instrument ID
1.	Biosafety Cabinet	Ascension, India	RRS/INS/CB/01
2.	CO ₂ Incubator	NUAIRE, USA	RRS/INS/CB/02
3.	Inverted tissue culture microscope	Nikon, Japan	RRS/INS/CB/08
4.	Automated micro plate reader	Biotek, USA	RRS/INS/MB/05

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7. METHOD

7.1. Outline of the method

The *in vitro* cytotoxicity was performed for the test substance on Human Keratinocyte (HaCaT) cell line to find a toxic concentration of the test substance by MTT assay. The percentage cytoprotective activity against UV induced damage was examined at nontoxic concentration.

7.2 Cell line and Culture medium:

HaCaT (Human Keratinocytes cells) was procured from AddexBio, U.S. Stock cells were cultured in DMEM-HG supplemented with 10% inactivated Fetal Bovine Serum (FBS), penicillin (100 IU/mL), streptomycin (100μg/mL) and amphotericin B (5μg/mL) in a humidified atmosphere of 5% CO₂ at 37°C until confluent. The cells were dissociated with TPVG solution (0.2% trypsin, 0.02% EDTA, 0.05% glucose in PBS). The stock cultures were grown in 25 cm² culture flasks and all experiments were carried out in 96 microtitre plates (Tarsons India Pvt. Ltd., Kolkata, India).

7.3 Preparation of test solution

10mg of test product was weighed and dissolved in DMEM-HG medium supplemented with 2% inactivated FBS to obtain a stock solution of 10mg/mL. Furthermore, serial-two fold dilutions were prepared from the stock solution to prepare lower concentrations for cytotoxicity testing.

10mg of test product was weighed and dissolved in DMEM-HG medium supplemented with 2% inactivated FBS to obtain a stock solution of 10mg/mL. Furthermore, non-toxic concentrations were prepared from the stock solution for further studies.

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7.4. Cytoprotective activity

Initially, the cytotoxicity study in HaCaT cells was carried out by standard methods to determine CTC50 value. 1.5 x 10⁵ cells were seeded in 6 well plate after reaching 70-80% confluency, the study was initiated. The supernatants were aspirated from wells, cultures were washed with DPBS and 2mL of DPBS was added to each well. To perform the Irradiation part of the assay, the cells were irradiated at room temperature for 50 minutes through the lid with 1.7mW/cm² (5 J/cm²). Non Irradiated control was kept at room temperature in a dark box for approximately 50 minutes. After irradiation, DPBS was decanted and the cells were washed once with DPBS. The DPBS was replaced with culture medium containing test substances and the plates were incubated at 37° ± 1°C for 24 hours. The plates were then incubated at 37° C for 24 h in 5% CO₂ atmosphere, and microscopic examination was carried out and observations were noted after 24h, the test solutions in the wells were discarded and MTT was added with DPBS to each well. The plates were gently shaken and incubated for 3 h at 37° C in 5% CO₂ atmosphere. The supernatant was removed and Isopropanol was added and the plates were gently shaken to solubilize the formed formazan. The absorbance was measured using a microplate reader at a wavelength of 570 nm.

8. RESULTS

Table 1: *In vitro* cytoprotective activity of test product in Human Keratinocytes (HaCaT) cell line against UV induced damage.

Sl. No.	Samples	Concentration tested	% Protection
1.	Tocyen Beauty Cream	1000µg/mL	67.47 ± 0.09
		500μg/mL	32.45 ± 0.08

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9. DISCUSSION AND CONCLUSION

The test product Tocyen Beauty Cream tested for *in vitro* cytotoxicity studies against Human Keratinocyte cell line by MTT assay exposing the cells to different concentrations. The CTC₅₀ value of the test product on HaCaT cell line was above 1000μg/mL. The *in vitro* cytoprotective activity of test product was evaluated in Human Keratinocyte (HaCaT) cell line at non-toxic concentrations of the test product. When the cells were subjected to UV and treated with the test product, the percentage protection exhibited by the test product against the control, Tocyen Beauty Cream (RR222098) showed the protection of 67.47 ± 0.09% and 32.45 ± 0.08% at 1000μg/mL and 500μg/mL respectively.

10. ARCHIVING

- Test Samples will be stored for 3 months after the final report submission.
- Raw data, documents, report will be archived for 3 years.

11. REPORT DISTRIBUTION

• Sponsor : One signed final report (Copy no. 1/2) in original.

• Archives : One signed final report (Copy no. 2/2) in original along with raw data file.

*****End of the report****