

1st International Congress on Analytical Chemistry, Electrochemistry and Separation Techniques



October 15th-16th, 2022

CYTOTOXICITY AND ANTI-INFLAMMATORY EFFECT OF LICHEN EXTRACT ON HACAT CELLS

Maya Abir TARTOUGA¹, Ibtissem ZEGHINA¹, Mohamed Badreddine MOKHTARI¹, Stefania MARZOCCO², Ibtissem ELOUAR¹

¹ Department of Animal Biology, Faculty of Natural and Life Sciences, University Frères Mentouri Constantine 1, route Ain el bey 25000, Constantine Algeria.

² Department of Pharmacy, School of Pharmacy, University of Salerno, Via Giovanni Paolo II 132, I-84084 Fisciano, Italy.

Email*: Tartouga.abirmaya@umc.edu.dz

INTRODUCTION:

Lichens are symbiotic organisms associating fungi with algae or cyanobacteria, they are used in traditional medicine to treat different diseases such as pulmonary diseases, wounds, and arthritis. Their secondary metabolites are known for their potent biological activities. The aim of this study is to evaluate the effect of lichen alcoholic extract on HaCaT cells viability and its anti-inflammatory activity through the inhibition of ROS and TNF-α produced by LPS stimulated HaCaT cells.

Lichen extraction Lichen powder Extraction with alcohol at 40 °C extract

Fig. 1. Lichen extraction

Total phenolic and flavonoids contents

Phenolic and flavonoids compounds of the alcoholic lichen extract were determined using a 96 well plates [1]; [2].

Cell culture

HaCaT cells were cultured, at 37 C° and 5% CO2,in Dulbecco's modified Eagle's medium (DMEM) high glucose. 10% fetal bovine serum (FBS), 1% Penicillin (100 U/mL) and streptomycin (100 mg/mL).

Cell viability assay

The effect of the extract on cell viability was assessed by the MTT assay as previously reported [3].

REFRENCES

- [1] L.Müller, S.Gnoyke, A.Popken, V. Böhm. LWT Food Sci Technol .2010; 43:992–9.
- [2] G.Topçu, A.Bilici, C. Sarikürkcü, M. Öztürk, A. Ulubelen. *Food Chem.* **2007**;103:816–22.
- [3] M. Alilou, S. Marzocco, D. Hofer, SF. Rapa, et al. J Nat Prod. 2020;83:245
- [4] SF.Rapa, B.Waltenberger, P. Simona, R. Siracusa, et al. *The FASEB J.* **2020**:1576–90.

Intracellular reactive oxygen release (ROS) production

Intercellular ROS formation was measured using the H2DCF-DA probe (2',7'-dichlorofluoresceindiacetate) as depicted [4].

TNF-α release

TNF- α levels were determined using the Diaclone human TNF- α Elisa kit following the manufacturer's instructions. Results were expressed as pg/mL.

RESULTS

Table 1. Total phenolic and flavonoids contents in lichen extract

	TPC (µg GAE/ml)	TFC (µg QE/ml)
Extract	++++	++

Cell viability:

The extract was not toxic on HaCaT cells, where the viability was 100% (Data not shown).

ROS release

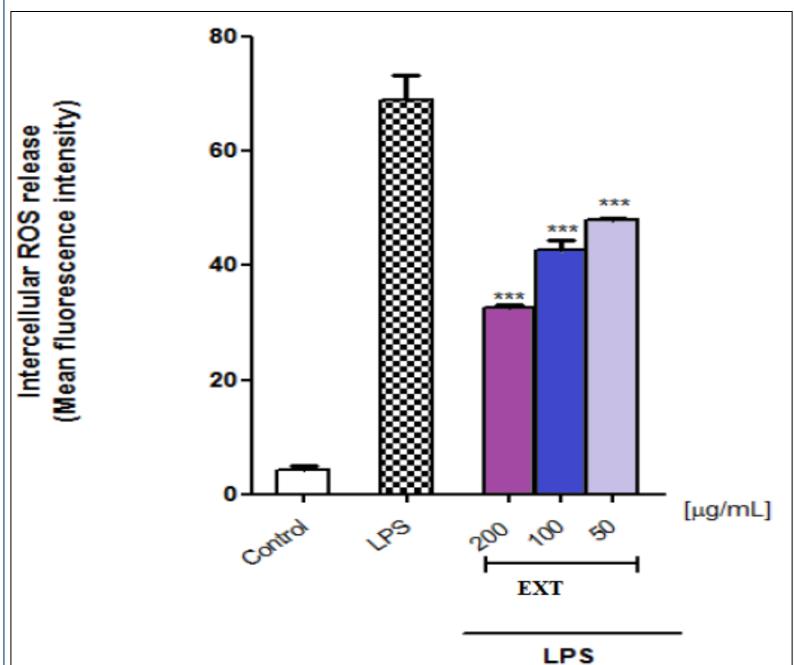


Fig. 2. Effect of lichen extract on the intracellular ROS levels In LPS stimulated HaCaT cells.

TNF-α release

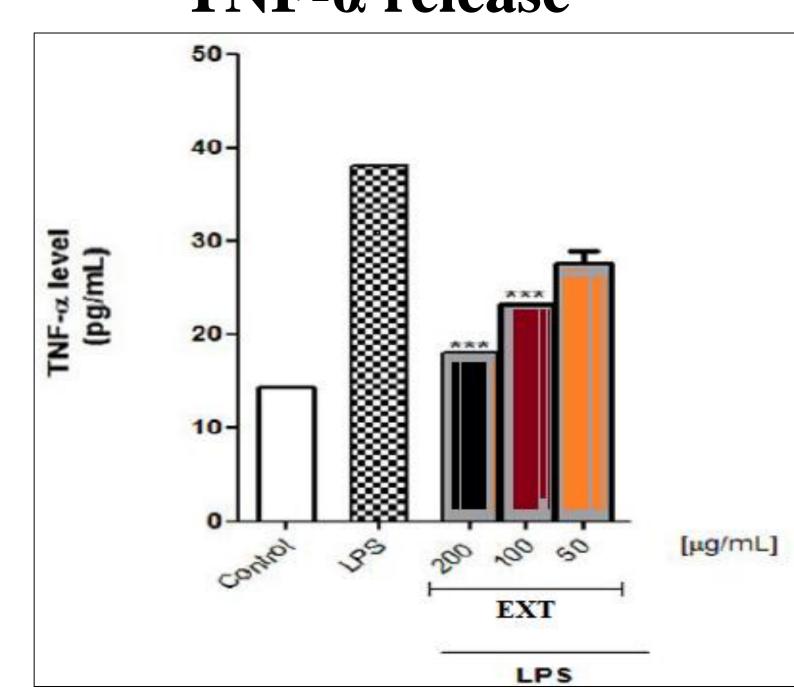


Fig. 4. Effect of the *extract* on TNF-α levels (pg/mL) in LPS Stimulated HaCaT cells.

CONCLUSION

The lichen extract could be a potent natural safe source of bioactive compounds with anti-inflammatory effects.