

[Published: 17 May 2021](#)

# Nerolidol, a sesquiterpene, attenuates oxidative stress and inflammation in acetic acid-induced colitis in rats

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*Molecular and Cellular Biochemistry* **476**, 3497–3512 (2021)

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## Abstract

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Targeting oxidative stress and inflammation by novel dietary compounds of natural origin convincingly appears to be one of the most important therapeutic strategies to keep inflammatory bowel diseases (IBD) such as ulcerative colitis disease in remission. It is imperative to investigate naturally occurring plant-derived dietary phytochemicals that are receiving attention for their therapeutic benefits to overcome the debilitating conditions of IBD. In the present study, the effect of nerolidol (NRD), a monocyclic sesquiterpene found in German Chamomile tea, was investigated in acetic acid-induced colitis model in Wistar rats. NRD was orally administered at a dose of 50 mg/kg/day either for 3 days before or 30 min after induction of IBD for 7 days, after intrarectal administration of acetic acid. The body weight, macroscopic, and microscopic analyses of the colon in different experimental

groups were observed on days 0, 2, 4, and 7. Acetic acid caused significant reduction in body weight and induced macroscopic and microscopic ulcer along with a significant decline of antioxidants, concomitant to increased malondialdehyde (MDA), a marker of lipid peroxidation, and myeloperoxidase (MPO) activity, a marker of neutrophil activation. Treatment with NRD significantly improved IBD-induced reduction in body weight, improved histology, inhibited MDA formation, and restored antioxidants along with reduced MPO activity. Acetic acid also induced the release of pro-inflammatory cytokines and increased calprotectin, released by neutrophils under inflammatory conditions. NRD treatment significantly reduced calprotectin and pro-inflammatory cytokines. NRD treatment showed potential to improve disease activity and inhibit oxidative stress, lipid peroxidation, and inflammation along with histological preservation of the colon tissues.

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## Data availability

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The manuscript data are well archived with the principal investigator, Prof. Salim MA Bastaki. The manuscript data will be available to the editors upon reasonable request from Prof. Salim Bastaki.

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Contributions

SMAB and SO have conceptualized the study.

SMAB designed the experiments, and interpreted the data.

NA has performed test treatments, animal care,

experiments, and biochemical estimations. NA carried out

the statistical analysis of all the data collected and archived

all the data. EA performed the histopathological studies

and interpreted the observations. SMBA and SO wrote the

first draft of manuscript and SMBA significantly edited the

manuscript. SO and SMBA have revised and submitted the

final manuscript. All authors read and approved the final

manuscript.

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Ethics declarations

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Competing interest

The authors are grateful for the financial support from the

United Arab Emirates University and for providing

facilities to conduct the experiments. The authors declare

that they have no competing interests.

Additional information

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## About this article

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### Cite this article

Bastaki, S.M.A., Amir, N., Adeghate, E. *et al.* Nerolidol, a sesquiterpene, attenuates oxidative stress and inflammation in acetic acid-induced colitis in rats. *Mol Cell Biochem* **476**, 3497–3512 (2021). <https://doi.org/10.1007/s11010-021-04094-5>

Received	Accepted	Published
05 September 2020	29 January 2021	17 May 2021

### Issue Date

September 2021

### DOI

<https://doi.org/10.1007/s11010-021-04094-5>

### Keywords

**Acetic acid**   **Colitis**   **Inflammation**   **Nerolidol**

**Oxidative stress**   **Phytochemicals**   **Rats**



