

# Anti-hyperglycemic and hepato-protective effects of thymoquinone (Nigella sativa oil) in diabetes

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## INTRODUCTION

Thymoquinone is the major component of Nigella sativa volatile oil, with antioxidant, antitumoral and anti-inflammatory properties. We explored the glycemetic and hepatic effects of thymoquinone in streptozotocin-treated rats.

## MATERIAL & METHODS

Thymoquinone was administrated to streptozotocin-induced diabetic female rats (25mg /kg/day) during 21 days.

Glycemic fluctuation was scrutinized and hepatic effects were assessed through biochemical analysis.



## RESULTS & DISCUSSION

Thymoquinone significantly reduced (-26%) the hyperglycemia induced by Nicotinamide/Streptozocin.

In vitro enzymatic evaluation showed that thymoquinone inhibited the  $\alpha$ -glucosidase with an estimated half maximal inhibitory concentration (IC<sub>50</sub>) = 125.03  $\mu$ g / ml

This hypoglycemic effect seems strong with a Thymoquinone IC<sub>50</sub> much lower than the IC<sub>50</sub> of the reference hypoglycemic drug, acarbose = 388.8  $\mu$ g / ml )

Biologically, thymoquinone improved the hepatic markers in treated diabetic rats :

ALT: 39.9  $\pm$  2 IU / L, GGT: 1.2  $\pm$  0.5 IU / L, Total bilirubin: 0.045  $\pm$  0.01 mg / dL, alkaline phosphatase 305.7  $\pm$  83 IU /L versus non-diabetic rats:

ALT: 22.1  $\pm$  2, GGT: 0.87  $\pm$  0.1 IU / L, Total bilirubin: 0.023  $\pm$  0.004 mg / dl, alkaline phosphatase 430  $\pm$  75 IU /L

## CONCLUSION

Thymoquinone significantly reduced blood sugar in diabetic murine models and acts as an oral anti-diabetic drug: its anti-hyperglycemic action is partly explained by a potent  $\alpha$ -glucosidase inhibition.

Thymoquinone might also have positive effects on the hepatic biological and histological parameters.