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# Antioxidant Activity of Methanolic Extract of Flowers of Nerium Oleander Against Ccl4-induced Liver Injury in Rats

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### **Abstract**

To investigate the inhibitor and hepatoprotective activity of methanolic flower extract of poisonous plant against CCl4-induced hepatotoxicity in rats. In vitro inhibitor activity of methanolic extract of flowers of poisonous plant (MENO–F) was evaluated by varied assays, together with reducing power, supermolecule peroxidation, DPPH, ABTS, superoxide, chemical group radicals and metal chelation.

# **Keywords**

inhibitor • hepatoprotective • alkalic enzyme • haematoidin • hepatotoxicity

# **About the Study**

The hepatoprotective and in vivo inhibitor activity of MENO-F were evaluated against CCl4—induced internal organ harm in rats. The MENO-F at dose of a hundred, two hundred and four hundred mg/kg were administered orally once daily for seven days. liquid body substance catalyst levels of serum salt salt aminopherase (AST), humour salt pyruvate aminopherase (ALT), humour alkalic enzyme (ALP) and total haematoidin were calculable beside estimation of SOD (SOD) and malondialdehyde (MDA) levels in liver tissues. any histopathological examination of the liver sections was dispensed to support the induction of hepatotoxicity and hepatoprotective effectuality.

The extract showed potent activities on reducing power, lipide peroxide, DPPH, ABTS, superoxide, chemical group and metal chelation. The considerably elevated body fluid catalyst levels of AST, ALT, mountain and total haematoidin were found to be reconditioned towards standardisation considerably by the MENO-F in a very dose dependent manner with most hepatoprotection at four hundred mg/kg dose level. The histopathological observations supported the organic chemistry evidences of hepatoprotection. Elevated level of SOD and shrunken level of MDA any strengthen the hepatoprotective observations.

## Conclusion

The results of this study powerfully reveal that MENO-F has potent inhibitor activity and hepatoprotective activity against CCl4-induced internal organ harm in experimental animals.

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