

MANAGEMENT OF HYPERLIPIDEMIA IN ANIMAL MODEL WITH MORINGA OLEIFERA

**Dr. Sana Imran¹, Dr. Syeda Afroz², Dr. Hafiza Tuseef Sayyar³, Dr. Abdullah⁴, Dr. Nabiha Iqbal⁵,
Dr. Aniq Naz⁶, Wasima Anjum⁷ Azmat Ara⁸**

¹ Assistant Professor, Department of Pharmacology, Jinnah Sindh Medical University, Karachi, Pakistan.

² Associate Professor, Department of Pharmacology, Faculty of Pharmacy and Pharmaceutical Sciences, University of Karachi, Karachi, Pakistan

³ Assistant Professor, Department of Pharmacology, Bahria University of Health Sciences, Karachi, Pakistan.

⁴ Lecturer, Department of Pharmacology, Faculty of Pharmacy and Health Sciences, University of Baluchistan, Quetta, Pakistan

⁵ Assistant professor, Department of Pharmacology, Bright international College, Lahore, Pakistan.

⁶ Research Scholar, Department of Pharmacology, Faculty of Pharmacy and Pharmaceutical Sciences, University of Karachi, Karachi, Pakistan

⁷ Department of Chemistry, Faculty of Science, University of Karachi, Karachi, Pakistan

⁸ Research Scholar, Department of Pharmaceutics, Faculty of Pharmacy and Pharmaceutical Sciences, University of Karachi, Karachi, Pakistan

Corresponding Author: Dr. Sana Imran

Assistant Professor, Department of Pharmacology, Jinnah Sindh Medical University, Karachi, Pakistan.

ABSTRACT

Introduction:

Hyperlipidemia, characterized by elevated cholesterol and triglyceride levels, poses a significant health risk, necessitating effective interventions. This study explores the potential of *Moringa oleifera*, a natural plant product rich in nutrients, as a promising candidate for regulating lipid profiles. *Moringa oleifera* has gained recognition for its diverse health benefits, making it an intriguing subject for managing hyperlipidemia.

Methodology:

Conducted at the Pharmacology department, University of Karachi, the study involved eighteen male Wistar rats distributed into three groups: a control group, a hyperlipidemic group treated with *Moringa oleifera*, and a hyperlipidemic group treated with the standard drug, Atorvastatin. Hyperlipidemia was induced through subcutaneous Dexamethasone injection for seven days. Blood samples confirmed hyperlipidemia, and subsequent treatment with *Moringa oleifera* leaf extract (0.06 mg) and Atorvastatin (0.6 mg) was administered for 21 days. Statistical analysis using SPSS 20 assessed lipid levels.

Results:

Both *Moringa oleifera* and Atorvastatin demonstrated significant reductions in total cholesterol, triglycerides, and LDL-C levels. Additionally, positive trends were observed in HDL-C levels. Statistical analysis revealed noteworthy changes in lipid levels from Day 0 to Day 21 in both treatment groups.

Conclusion:

The research highlights that *Moringa oleifera* proves to be a healthy and cost effective natural substitute to Atorvastatin, a standard hypolipidemia

Keywords: Hyperlipidemia, *Moringa Oleifera*, Atorvastatin

INTRODUCTION

Hyperlipidemia (Hyper-LP) is a state labelled as an increment in total cholesterol, triglycerides, Low Density Lipoprotein-Cholesterol (LDL-C), and a decrement in High Density Lipoprotein-Cholesterol (HDL-C) 1. The primary treatment is improving dietary and daily routines, trailed by synthetic & natural hypolipidemic drugs 2.

Moringa oleifera is one of the natural plant product cultivated in subtropical and tropical regions around the world 3. Leaves are most consuming part containing a variety of healthy nutrients such as amino acids, beta-carotene, vitamins and minerals 4. This natural plant product is beneficial in multiple health issues like deranged lipid profile, diabetes mellitus, bleeding disorders, GI disturbances 5. Thus, *Moringa oleifera* acts as a functional food.

According to the Functional Food Science in Europe (FUFOSE), functional food is a food that has one or more beneficial effects in the body beyond basic nutrition that have an impact on health and reduce the risk of diseases 6.

The most prescribed alone or in combination synthetic hypolipidemic drugs are Statins. Atorvastatin is one of its example that effects the conversion of HMG-CoA to Mevalonic acid by impeding rate restraining cholesterol biosynthesis enzyme HMG-CoA reductase and lowers Low Density Lipoprotein-Cholesterol (LDL-C) 7.

METHOD

This research work was performed in the Pharmacology department, University of Karachi after the confirmation of ethical committee of ASRB (ASRB No/ 05848 /Pharm).

In this 3-week study, male healthy Wistar rats total eighteen in number were consumed. The rats were equally distributed into three research groups. Group-1 was Control (Non-Hyperlipidemic) group, Group-2 was Hyperlipidemic – *Moringa oleifera* treated group & Group-3 was Hyperlipidemic – Atorvastatin treated group.

Hyperlipidemia was produced in Group-2 and Group-3 Wistar rats by injecting Dexamethasone subcutaneously for seven days 8.

Afterwards hyperlipidemia was confirmed from blood samples gathered from each group. The *Moringa oleifera*'s dose of 0.06 mg was given in Group-2 rats, and the Atorvastatin's dose of 0.6 mg was given in Group-3 rats orally per day in the morning for 21 days research work. Both the drugs were given after the confirmation of hyperlipidemia considering Day-0 till Day- 21 labelling last day of research.

The objective of this study was to determine the lipid lowering effects in induced Wistar rats by giving leaf extract of *Moringa oleifera* that seems to be an alternative source to regularize body's lipid profile,

The p-value lesser than 0.05 was considered noteworthy in all statistical analysis. The data feeding and analysis was done on SPSS 20.

RESULTS

The study work done on both Group – 2 and Group – 3 revealed that *Moringa oleifera* has comparable reduction effects on total cholesterol, triglyceride and LDL-C levels. Similarly, *Moringa oleifera* has matchable healthy effect on HDL-C level seen from Day – 0 to Day – 21 as exhibited in Tables-1, 2, 3 & 4.

TABLE-1: COMPARISON IN MEAN TOTAL CHOLESTEROL (mg/dL) LEVEL AMONG DIFFERENT EXPERIMENTAL GROUPS FROM DAY 0 - DAY 21

GROUPS	DAY – 0	DAY - 21	p-VALUE
Group – 1	158.50 ± 2.12	172.50 ± 3.70	0.0087*
Group – 2	208.50 ± 3.54	189.50 ± 7.33	0.02911*
Group – 3	212.00 ± 2.83	195.00 ± 2.16	0.0011*

TABLE-2: COMPARISON IN MEAN TRIGLYCERIDE (mg/dL) LEVEL AMONG DIFFERENT EXPERIMENTAL GROUPS FROM DAY 0 - DAY 21

GROUPS	DAY – 0	DAY – 21	p-VALUE
Group – 1	102.50 ± 10.61	102.00 ± 8.83	0.9535
Group – 2	137.50 ± 6.36	111.75 ± 1.71	0.0011*
Group – 3	140.00 ± 5.66	113.75 ± 1.71	0.0007*

TABLE-3: COMPARISON IN MEAN HDL-C (mg/dL) LEVEL AMONG DIFFERENT EXPERIMENTAL GROUPS FROM DAY 0 - DAY 21

GROUPS	DAY – 0	DAY – 21	p-VALUE
Group – 1	60.00 ± 2.83	59.50 ± 2.65	0.8407
Group – 2	50.00 ± 1.41	58.50 ± 1.29	0.0018*
Group – 3	49.00 ± 1.41	57.50 ± 1.29	0.0018*

TABLE-4: COMPARISON IN MEAN LDL-C (mg/dL) LEVEL AMONG DIFFERENT EXPERIMENTAL GROUPS FROM DAY 0 - DAY 21

GROUPS	DAY – 0	DAY - 21	p-VALUE
Group – 1	107.50 ± 3.54	113.25 ± 2.99	0.1014
Group – 2	143.50 ± 2.12	131.75 ± 2.75	0.0065*
Group – 3	151.00 ± 2.83	135.75 ± 2.22	0.0018*

Values are expressed in Mean ± S.D

S.D = Standard Deviation

* = Significant

GROUP 1: Control Group (Non-Diabetic, Non-Hyper-LP)

GROUP 2: Moringa oleifera treated Hyper-LP group

GROUP 3: Atorvastatin treated Hyper-LP group

DISCUSSION

Our research work led from Day-0 till Day-21 explored *Moringa oleifera*'s healthy comparable effects on hyperlipidemia induced rats.

The research of Ghasi et al (2000), 9 represented the medicinal effect of MO in normalizing mean total cholesterol, triglyceride, HDL-C and LDL-C levels that in accord to our *Moringa oleifera* treated Group – 2 and suggestive of useful substitute of hyperlipidemia therapy

Atorvastatin treated Group – 3 showed that Atorvastatin was effective in the treatment of hyperlipidemia when taken from Day 0 to Day-21.

The research work performed by Lin, LY. et al (2014), 10 is consistent with our study showing Atorvastatin is a standard hypolipidemic drug that ameliorates hyperlipidemia in short-term study.

CONCLUSION

Moringa oleifera emerges as a promising, cost-effective, and natural alternative to Atorvastatin in regulating lipid profiles. The study provides valuable insights into the comparative efficacy of *Moringa oleifera* and Atorvastatin in a controlled experimental setting. These findings underscore the potential of *Moringa oleifera* as a dietary intervention for managing hyperlipidemia, encouraging further exploration of natural plant products as viable alternatives to synthetic drugs for lipid-related disorders.

REFERENCES

1. Vafa, M., E. Haghghatjoo and A. Ziaee. Effect of Apple Consumption on Lipid Profile of Hyperlipidemic and Overweight Men. *International Journal of Preventive Medicine*. 2011; 2 (2): 84-100.
2. Henley, E., L. Chang and S. Hollander. 2002. Treatment of Hyperlipidemia. *The Journal of Family Practice* 2002; 51(4):370-376.
3. Leone, A., Spada, A., Battezzati, A., et al. Cultivation, genetic, ethnopharmacology, phytochemistry and pharmacology of *Moringa oleifera* Leaves: An overview. *Int. J. Mol. Sci*. 2015; 16:12791–835.
4. Falowo, A.B., Mukumbo, F.E., Idamokoro, E.M., et al. Multi-functional application of *Moringa oleifera* Lam. in nutrition and animal food products: A review. *Food Res. Int*. 2018; 106:317–34.

5. Lalas S, Tsaknis J. Extraction and identification of natural antioxidants from the seeds of *Moringa oleifera* tree variety of Malawi. *J Am Oil Chem Soc.* 2002; 79: 677–683.
6. Anonymous. 2010. *Functional Food*. Luxembourg: Publications Office of the European Union.
7. Antoniades C, Bakogiannis C, Leeson P, et al. Rapid, direct effects of statin treatment on arterial redox state and nitric oxide bioavailability in human atherosclerosis via tetrahydrobiopterin-mediated endothelial nitric oxide synthase coupling. *Circulation* 2011; 124(3): 335-45.
8. Kumar VR, Inamdar MN, Nayeemunnisa, Viswanatha GL. Protective effect of lemongrass oil against dexamethasone induced hyperlipidemia in rats: possible role of decreased lecithin cholesterol acetyl transferase activity. *Asian Pac J Trop Med.* 2011; 4: 658-60. [DOI:10.1016/S1995-7645(11)60167-3]
9. Ghasi S, Nwobodo E, Ofili J. Hypocholesterolemic effects of crude extract of leaf of *Moringa oleifera* lam in high-fat diet fed Wistar rats. *J Ethnopharmacol.* 2000;69(1):21–5.
10. Lin, LY., Huang, CC., Chen, JS. et al. Effects of pitavastatin versus atorvastatin on the peripheral endothelial progenitor cells and vascular endothelial growth factor in high-risk patients: a pilot prospective, double-blind, randomized study. *Cardiovasc Diabetol* 13, 111 (2014). <https://doi.org/10.1186/s12933-014-0111-1>