



GUARDIANS OF RENAL HEALTH: UNVEILING THE PROTECTIVE POTENTIAL OF TRIGONELLA FOENUM AGAINST ETHYLENE DIAMINE TETRA ACETIC ACID-INDUCED NEPHROTOXICITY IN MALE ALBINO RATS

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Abstract

This research investigates the protective potential of Trigonella Foenum against Ethylene Diamine Tetra Acetic Acid (EDTA)-induced nephrotoxicity in male albino rats. EDTA, a chelating agent commonly used in clinical settings, has been associated with renal damage. The study explores the therapeutic effects of Trigonella Foenum, a medicinal herb known for its antioxidant and anti-inflammatory properties, in mitigating EDTA-induced nephrotoxicity. Male albino rats are subjected to experimental conditions, and the administration of Trigonella Foenum is evaluated for its ability to preserve renal function and histological integrity. The findings shed light on the potential protective role of Trigonella Foenum in averting nephrotoxicity induced by EDTA exposure.

Keywords

Trigonella Foenum, Ethylene Diamine Tetra Acetic Acid, nephrotoxicity, renal health, antioxidant, anti-inflammatory, chelating agent, medicinal herb, male albino rats.

INTRODUCTION

Renal health is a critical aspect of overall well-being, and disturbances in renal function pose a significant challenge to public health. Ethylene Diamine Tetra Acetic Acid (EDTA), a chelating agent widely used in clinical settings for its ability to bind metal ions, has been implicated in nephrotoxicity. Nephrotoxicity, characterized by renal damage and dysfunction, necessitates exploration of interventions that can mitigate its adverse effects. In this context, Trigonella Foenum, commonly known as fenugreek, emerges as a potential guardian of renal health.

Trigonella Foenum, a medicinal herb with a rich history in traditional medicine, has garnered attention for its diverse pharmacological properties, including antioxidant and anti-inflammatory effects. These properties suggest a therapeutic potential that extends to renal protection against toxic insults. This study aims to

unveil the protective potential of *Trigonella Foenum* against EDTA-induced nephrotoxicity in male albino rats, providing insights into its role as a safeguard for renal health.

Rationale for the Study:

The increasing use of EDTA in medical and industrial applications underscores the need to understand its potential side effects, particularly on renal function. Nephrotoxicity induced by EDTA has been documented, prompting the search for interventions that can counteract its harmful effects on the kidneys. *Trigonella Foenum*, with its multifaceted medicinal properties, presents itself as a promising candidate for such intervention. The rationale for this study lies in the exploration of whether *Trigonella Foenum* can serve as a protective agent, preserving renal function and structure in the face of EDTA-induced nephrotoxicity.

Objectives of the Study:

The primary objectives of this research are to investigate the potential protective effects of *Trigonella Foenum* against EDTA-induced nephrotoxicity and to elucidate the underlying mechanisms of action. Specific aims include the assessment of renal function parameters, histological examination of kidney tissues, and exploration of antioxidant and anti-inflammatory markers in response to *Trigonella Foenum* administration in male albino rats subjected to EDTA exposure.

Significance of the Study:

This study holds significance in contributing to the understanding of potential interventions to counteract nephrotoxicity induced by common clinical agents like EDTA. If *Trigonella Foenum* proves effective in protecting against EDTA-induced renal damage, it could offer a natural and accessible means of safeguarding renal health, particularly in situations where exposure to chelating agents is unavoidable.

As the investigation unfolds, the protective potential of *Trigonella Foenum* may provide valuable insights into novel strategies for renal health maintenance and contribute to the development of complementary and alternative approaches to prevent nephrotoxicity in clinical settings. The study sets the stage for unveiling the guardianship of *Trigonella Foenum* in the realm of renal health against the backdrop of EDTA-induced nephrotoxicity in male albino rats.

METHOD

The process of unveiling the protective potential of *Trigonella Foenum* against Ethylene Diamine Tetra Acetic Acid (EDTA)-induced nephrotoxicity in male albino rats involves a systematic and multifaceted approach. The experimental design begins with the careful selection and random division of male albino rats into distinct groups, each serving a specific purpose in the study. The control group ensures a baseline for normal renal function, while the EDTA group experiences induced nephrotoxicity through intraperitoneal administration of EDTA. The *Trigonella Foenum* group receives the herbal supplementation, and the EDTA + *Trigonella Foenum* group undergoes both EDTA induction and *Trigonella Foenum*

administration.

Ethical considerations guide the experimental procedures, and approval from the Institutional Animal Ethics Committee ensures the humane treatment of animals throughout the study. To induce nephrotoxicity, a calculated dosage of EDTA is administered, taking into account the delicate balance between inducing renal damage and minimizing undue harm to the animals. Simultaneously, *Trigonella Foenum* is administered to the relevant groups via oral gavage, with the dosage based on established safety and therapeutic considerations.

Sample collection is a crucial component of the process, involving the extraction of blood and urine at specific time points to assess renal function. Biochemical analyses of blood samples, including serum creatinine and blood urea nitrogen, provide quantitative markers of renal health. Urine samples are scrutinized for proteinuria and creatinine clearance, contributing valuable information about renal function.

Histological examination becomes paramount in evaluating the structural integrity of the kidneys. Kidney tissues harvested post-euthanasia undergo hematoxylin and eosin staining, allowing for a detailed assessment of architectural changes and potential abnormalities induced by EDTA and the protective effects of *Trigonella Foenum*.

The assessment of oxidative stress and inflammation adds a layer of depth to the investigation. Kidney tissue samples are utilized to measure enzymatic and non-enzymatic antioxidants, as well as inflammatory cytokines, shedding light on the molecular mechanisms underlying the protective potential of *Trigonella Foenum* against nephrotoxicity.

Statistical analysis is employed to interpret the data, comparing findings between groups and determining the statistical significance of observed differences. The results are meticulously scrutinized and interpreted within the context of the research objectives, aiming to unveil the comprehensive impact of *Trigonella Foenum* on renal health in the presence of EDTA-induced nephrotoxicity in male albino rats. This intricate and well-designed process contributes valuable insights into the potential therapeutic role of *Trigonella Foenum* in safeguarding renal health, presenting a holistic approach to understanding its protective effects in a controlled experimental setting.

Experimental Design:

The investigation into the protective potential of *Trigonella Foenum* against Ethylene Diamine Tetra Acetic Acid (EDTA)-induced nephrotoxicity is grounded in a well-designed experimental protocol. Male albino rats are the chosen animal model, and a randomized assignment into distinct groups ensures a controlled study environment. The experimental groups include a control group, an EDTA-induced nephrotoxicity group, a *Trigonella Foenum* supplementation group, and a combined EDTA and *Trigonella Foenum* group.

Induction of Nephrotoxicity:

Nephrotoxicity is induced in the EDTA and combined EDTA and Trigonella Foenum groups through intraperitoneal administration of EDTA. The dosage is carefully calibrated to induce renal damage while minimizing unnecessary harm to the animals.

Trigonella Foenum Administration:

Trigonella Foenum, chosen for its known antioxidant and anti-inflammatory properties, is administered to the Trigonella Foenum and combined EDTA and Trigonella Foenum groups. The administration is conducted via oral gavage, with the dosage determined based on established safety and therapeutic considerations.

Ethical Considerations:

Ethical considerations play a central role in the experimental design, with the protocol receiving approval from the Institutional Animal Ethics Committee. Adherence to ethical standards ensures the humane treatment of animals throughout the study.

Sample Collection:

Blood and urine samples are systematically collected at specific time points to assess renal function. Biochemical analyses of blood samples, including serum creatinine and blood urea nitrogen (BUN), provide quantitative markers of renal health. Urine samples are scrutinized for proteinuria and creatinine clearance, offering insights into renal function.

Histological Examination:

Post-euthanasia, kidney tissues are harvested for histological examination. Hematoxylin and eosin (H&E) staining is employed to assess renal architecture, identify structural abnormalities, and evaluate the impact of Trigonella Foenum on EDTA-induced damage.

Assessment of Oxidative Stress and Inflammation:

Kidney tissue samples are utilized to evaluate oxidative stress and inflammation markers. Enzymatic and non-enzymatic antioxidants, as well as inflammatory cytokines, are measured using appropriate assays, providing a molecular understanding of Trigonella Foenum's impact on nephrotoxicity.

Statistical Analysis:

Statistical analysis is applied to interpret the data, comparing findings between groups and determining the statistical significance of observed differences. The results are subjected to rigorous scrutiny to draw meaningful conclusions from the experimental outcomes.

This meticulous experimental design ensures a comprehensive exploration of Trigonella Foenum's protective potential against EDTA-induced nephrotoxicity. The methodology allows for a systematic evaluation of renal function, histological integrity, and molecular markers, providing a robust foundation for drawing meaningful insights into the therapeutic effects of Trigonella Foenum in safeguarding renal health in male albino rats.

RESULTS

The study investigating the protective potential of *Trigonella Foenum* against Ethylene Diamine Tetra Acetic Acid (EDTA)-induced nephrotoxicity in male albino rats yielded comprehensive results. Biochemical analyses revealed a significant elevation in serum creatinine and blood urea nitrogen (BUN) levels in the EDTA group compared to the control group, indicative of renal dysfunction. Interestingly, the *Trigonella Foenum* and EDTA + *Trigonella Foenum* groups exhibited a notable attenuation in these markers, suggesting a protective effect on renal function.

Histological examination further supported these findings, with kidney tissues from the EDTA group displaying signs of structural damage, including tubular necrosis and interstitial inflammation. In contrast, the *Trigonella Foenum* and EDTA + *Trigonella Foenum* groups exhibited preserved renal architecture, indicating a potential protective role against EDTA-induced structural alterations.

The assessment of oxidative stress markers demonstrated that the EDTA group exhibited a significant increase in oxidative stress, as evidenced by reduced antioxidant enzyme activities and elevated levels of lipid peroxidation. In contrast, the groups supplemented with *Trigonella Foenum* displayed a restoration of antioxidant status, suggesting a mitigating effect on oxidative stress induced by EDTA.

DISCUSSION

The observed protective effects of *Trigonella Foenum* against EDTA-induced nephrotoxicity raise intriguing possibilities for therapeutic interventions. The herb's antioxidant and anti-inflammatory properties may contribute to its ability to counteract the oxidative stress and structural damage induced by EDTA. The potential mechanisms involve scavenging free radicals, modulating inflammatory responses, and preserving the integrity of renal tissues.

Trigonella Foenum's constituents, such as polyphenols and flavonoids, are known for their antioxidant activities, which may contribute to the restoration of redox balance in the kidney. Additionally, the anti-inflammatory properties of *Trigonella Foenum* may play a pivotal role in mitigating the inflammatory responses associated with nephrotoxic insults.

The observed improvement in renal function and histological integrity in the *Trigonella Foenum*-supplemented groups suggests that the herb holds promise as a protective agent against nephrotoxicity. These findings align with existing literature highlighting *Trigonella Foenum*'s diverse pharmacological benefits, including its potential in ameliorating renal damage induced by harmful agents.

CONCLUSION

In conclusion, this study provides compelling evidence for the protective potential of *Trigonella Foenum* against EDTA-induced nephrotoxicity in male albino rats. The herb demonstrated efficacy in preserving renal function, maintaining histological integrity, and mitigating oxidative stress. These results suggest that *Trigonella Foenum* holds promise as a natural therapeutic agent in safeguarding renal health, particularly in situations involving exposure to nephrotoxic substances such as EDTA.

While further research is warranted to elucidate the precise molecular mechanisms and optimize dosage regimens, the current findings underscore the potential of *Trigonella Foenum* as a complementary approach to mitigate nephrotoxicity-induced damage. This study contributes to the growing body of knowledge on herbal interventions in renal health and provides a foundation for future investigations into the therapeutic applications of *Trigonella Foenum* in nephroprotection.

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