

# Therapeutic Role and Safety Profile of Corticosteroids in Renal Disorders

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## Abstract

Corticosteroids are extensively used in the management of several renal disorders due to their potent anti-inflammatory and immunosuppressive properties <sup>(1)</sup>. They remain a cornerstone of therapy in conditions such as glomerulonephritis, nephrotic syndrome, lupus nephritis, acute interstitial nephritis, and renal transplantation <sup>(7)</sup>. Although their clinical benefits are well established, long-term corticosteroid therapy is associated with a wide range of adverse effects, particularly in patients with chronic kidney disease <sup>(3)</sup>. This review summarizes evidence published after 2015 regarding the therapeutic role and safety profile of corticosteroids in renal disorders.

## Keywords

Corticosteroids; Renal Disorders; Glomerulonephritis; Nephrotic Syndrome; Safety Profile

## Introduction

Corticosteroids have been widely used in nephrology for decades because of their ability to suppress inflammatory pathways and modulate immune responses <sup>(7)</sup>. They are commonly prescribed in glomerular diseases, nephrotic syndrome, lupus nephritis, acute interstitial nephritis, and in immunosuppressive regimens following renal transplantation <sup>(8)</sup>. Despite their therapeutic benefits, corticosteroids are associated with several adverse effects, particularly when used for prolonged durations <sup>(3)</sup>.

## Objectives

1. To evaluate therapeutic effectiveness of corticosteroids in renal disorders.
2. To assess safety profile in renal patients.
3. To summarize recent clinical evidence.

## Methodology

A literature search was conducted using PubMed, Google Scholar, and Scopus for studies published between 2015 and 2025. Search terms included “corticosteroids,” “renal disorders,” and “glomerulonephritis.” Randomized

controlled trials, meta-analyses, and cohort studies were included.

## Discussion

Steroids remain effective in immune-mediated renal disorders such as IgA nephropathy and lupus nephritis <sup>(6)(7)</sup>. However, adverse effects such as infections, hyperglycemia, and osteoporosis limit long-term use <sup>(3)(8)</sup>. Recent evidence supports low-dose and combination regimens to reduce toxicity

## References

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<sup>(4)</sup>.

## Conclusion

Corticosteroids remain essential in selected renal disorders but require individualized dosing and careful monitoring <sup>(7)</sup>. Steroid-sparing strategies are promising for safer long-term management <sup>(4)</sup>

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## Review of Literature Table

<b>Author &amp; year</b>	<b>Study design</b>	<b>Population</b>	<b>Interventions</b>	<b>Key Outcomes</b>	<b>Conclusion</b>
<b>Jääskeläinen et al., 2023</b>	Systematic review & meta-analysis	Children with acute pyelonephritis	Corticosteroids + antibiotics	Reduced Renal scarring	Adjunctive steroids beneficial
<b>Arivazhagan et al., 2022</b>	Randomized controlled trial	Infection-related GN patients	Prednisolone vs placebo	No renal recovery benefit; ↑ adverse events	Steroids not beneficial in IRGN
<b>Del Vecchio et al., 2024</b>	Retrospective cohort	IgA nephropathy patients	Corticosteroid therapy	↓ Proteinuria; ↑ infections	Efficacy with safety concerns
<b>Zhang et al., 2024</b>	Meta-analysis	Progressive IgA nephropathy	Low-dose steroids + leflunomide	Improved outcomes; better safety	Combination preferred
<b>Gkiourtzis et al., 2023</b>	Meta-analysis	Pediatric UTIs	Corticosteroids + antibiotics	Reduced renal scarring	Steroids beneficial
<b>Radhakrishnan et al., 2019</b>	Observational study	Adult minimal Change disease	Prednisone	High remission; relapses	First-line therapy
<b>Praga et al., 2015</b>	Review article	FSGS patients	Steroid therapy	Variable response; toxicity	Careful selection needed

## List of Abbreviations

CKD – Chronic Kidney Disease

GN – Glomerulonephritis

IgAN – IgA Nephropathy

IRGN – Infection-Related  
Glomerulonephritis

RCT – Randomized Controlled Trial

KDIGO – Kidney Disease: Improving  
Global Outcomes

FSGS – Focal Segmental  
Glomerulosclerosis

UTI – Urinary Tract Infection

SAE – Serious Adverse Event

MMF – Mycophenolate Mofetil

