

# Overview of IgA Nephropathy

**Immunoglobulin A nephropathy (IgAN)**, also known as Berger's disease, is a leading cause of chronic kidney disease and the most common primary glomerulonephritis worldwide.<sup>1,2</sup> The majority of patients with IgAN develop **end-stage kidney disease (ESKD)** within their lifetime.<sup>3,4</sup>



## What Is IgAN?

- IgAN is a **progressive, autoimmune, chronic kidney disease** that is primarily characterized by mesangial immunoglobulin A (IgA) deposition<sup>1,2</sup>
- The etiologic pathogenesis of IgAN is multifactorial and can be explained by the **four-hit cascade**, the outcome of which is kidney injury that can lead to ESKD<sup>5,6</sup>



## Symptoms

- IgAN typically manifests in adults aged **20 to 40 years** and is often diagnosed incidentally<sup>7,8</sup>
- IgAN is a heterogenous disease with variable presentation, and patients frequently experience disease progression **despite being asymptomatic**<sup>1,8,9</sup>

### Symptoms may include:

- **Hematuria** (both macroscopic and microscopic)<sup>1</sup>
  - In some patients, macroscopic hematuria is preceded by an upper respiratory infection<sup>10</sup>
- **Proteinuria**<sup>1</sup>
- **Fatigue**<sup>11</sup>
- **Edema**<sup>11</sup>



## Who Is Affected?

- The annual global incidence of IgAN is **2.5 cases per 100,000 persons**; however, this varies by region and ethnicity<sup>12,13</sup>
- There is considerable heterogeneity in outcomes across different races and ethnicities; Asian populations may face **higher risk of severe disease** than White populations<sup>6,12</sup>



## Diagnosis and Prognosis

- IgAN is diagnosed with a renal biopsy showing **predominant IgA1 staining**<sup>1,14</sup>
- In observational studies, most patients with IgAN develop **ESKD within 10 to 15 years** of diagnosis<sup>3</sup>
  - Even in patients previously considered "low risk" (proteinuria <1 g/d), there is a substantial risk of kidney failure within 15 years of diagnosis<sup>3</sup>

## Optimized Supportive Care

According to current guidelines, the main focus of disease management is to reduce the risk of progression, beginning with optimization of supportive care.<sup>14</sup> The objective of optimized supportive care is to slow down progressive chronic kidney disease and reduce cardiovascular risk.<sup>14</sup>



Assess cardiovascular risk<sup>14</sup>



Control blood pressure<sup>14</sup>



Decrease proteinuria<sup>14</sup>



Dietary sodium restriction<sup>14</sup>



Smoking cessation<sup>14</sup>



Weight control and exercise<sup>14</sup>

## Patients at Risk of Disease Progression Despite Maximal Supportive Care

- A significant portion of patients remain at risk of progression despite optimized supportive care<sup>14,15</sup>
- According to guidelines, patients who have proteinuria >1 g/d despite optimized supportive care for ≥90 days may consider enrolling in a clinical trial<sup>14</sup>
- Immunosuppressive therapy may be considered with caution<sup>14</sup>

### References:

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