

# Atrasentan for the Treatment of IgA Nephropathy: Interim Results from the AFFINITY Study

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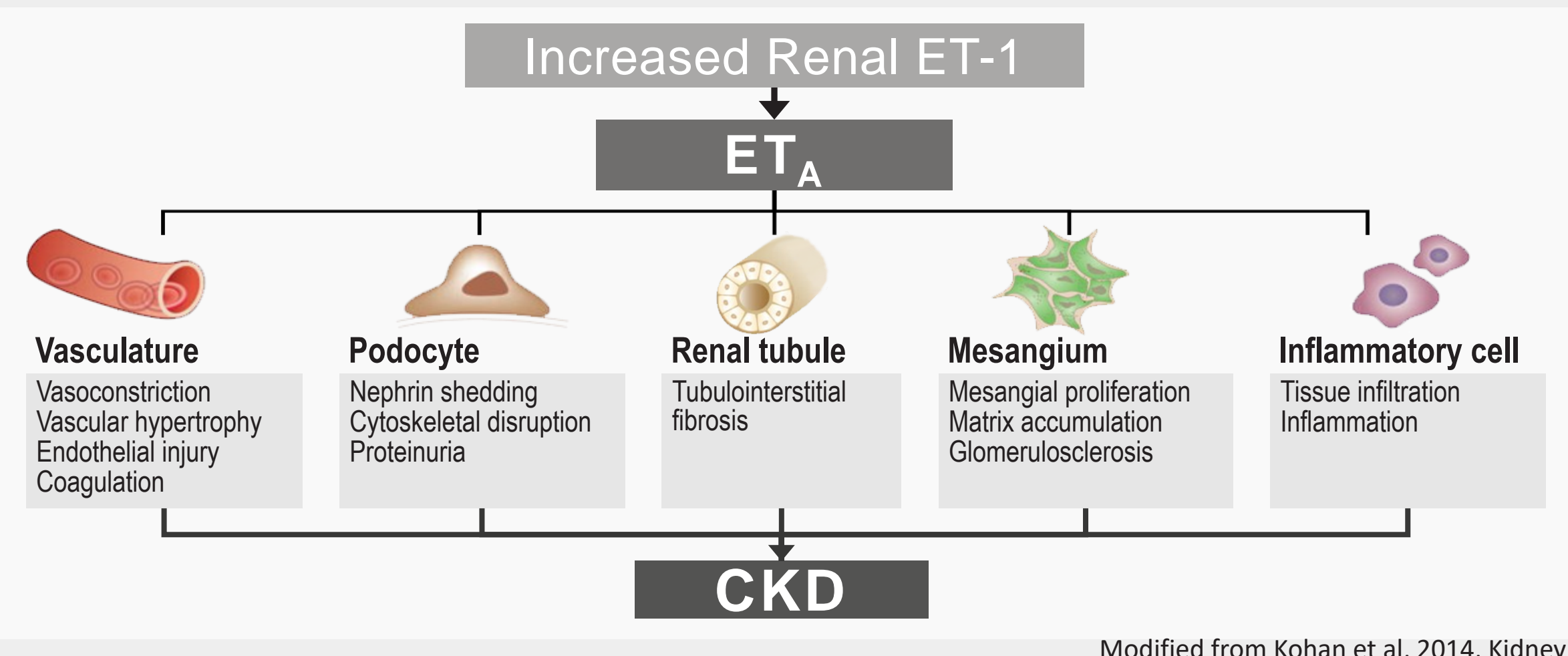
## Background/Methods

### IgA Nephropathy (IgAN)

- IgAN is the leading cause of primary glomerulonephritis, with a global incidence of 2.5 per 100,000 individuals per year<sup>1</sup>
- Approximately 30-45% of IgAN patients progress to end-stage kidney disease (ESKD) over a period of 20-25 years<sup>2-5</sup>
- Proteinuria is strongly associated with kidney disease progression in IgAN<sup>2,6-7</sup> and treatments that reduce proteinuria result in improved clinical outcomes in IgAN<sup>8-9</sup>

### Endothelin System Activation in IgAN

- Endothelin (ET-1) is a key contributor to progression of IgA nephropathy
- Elevated kidney ET-1 expression strongly & prospectively predicts progression of IgAN 12 months following kidney biopsy<sup>10</sup>
- Endothelin A (ET<sub>A</sub>) receptor activation drives mesangial cell activation, kidney inflammation & fibrosis, and proteinuria, all hallmarks of IgAN<sup>11-12</sup>
- Kidney ET-1 & ET<sub>A</sub> receptor levels are elevated in proteinuric patients with IgAN<sup>13-14</sup>



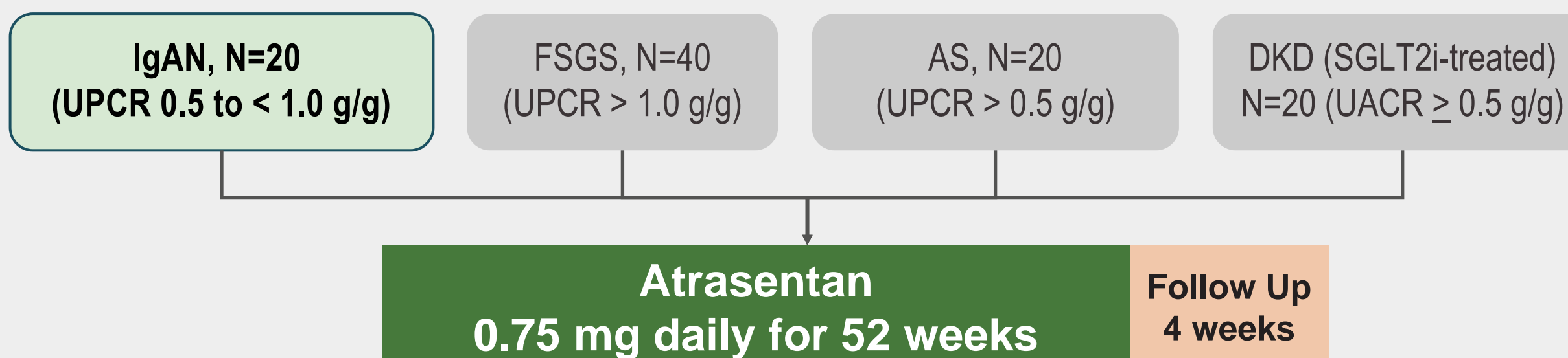
Modified from Kohan et al., 2014, Kidney Int

### Atrasentan\* has potential to treat IgAN patients at high risk of progression

- Atrasentan is a potent and highly-selective endothelin receptor A antagonist (K<sub>i</sub> = 0.034nM) with >1,800-fold selectivity over ETB (K<sub>i</sub> = 63.3nM)<sup>15</sup>
- Atrasentan has previously demonstrated clinically significant and sustained proteinuria reduction with an acceptable safety profile in over 5,100 patients with diabetic kidney disease (DKD)<sup>16-17</sup>
- In preclinical studies, atrasentan attenuates mesangial cell activation, glomerular and tubulointerstitial injury, and reduces proteinuria associated with IgAN<sup>18-21</sup>

### AFFINITY Study Design

- AFFINITY is a global, phase 2, open label basket study to assess the efficacy and safety of atrasentan in patients with proteinuric glomerular diseases (IgAN, focal segmental glomerulosclerosis [FSGS], Alport syndrome [AS], and DKD) at risk of progressive kidney function loss (NCT04573920)



#### Key Eligibility Criteria, IgAN Cohort

- Biopsy-proven IgAN
- Maximally-tolerated and optimized dose of a RAS inhibitor (RASi) for ≥ 12 weeks prior to screening
- UPCR of 0.5 to < 1.0 g/g (56.5 mg/mmol to <113 mg/mmol) based on first morning void urine collected at screening
- eGFR ≥ 30 mL/min/1.73 m<sup>2</sup>

#### Key Study Endpoints

- Change from baseline at week 12 in UPCR, based on average of two 24-hour collections
- Analysis based on an MMRM model of change from baseline in UPCR
- Adverse Event (AE) type, incidence, severity, seriousness and relatedness

## Baseline and Safety

### AFFINITY IgAN Cohort

- The AFFINITY IgAN cohort enrolled 20 patients with biopsy-confirmed IgAN
- All patients received concurrent, max-tolerated and optimized RASi at least 12 weeks prior to study and throughout the study period
- 70% of patients had baseline total urine protein >1 g/day despite optimized RASi treatment, representing an IgAN population at high risk for progression
- Mean treatment duration was 45 weeks (range 13-53 weeks) as of data cut-off October 19, 2022

DEMOGRAPHICS, N=20	
Age, years, median (Q1, Q3)	45 (35, 58)
Women, n (%)	10 (50)
Race, n (%), Asian	9 (45)
White	9 (45)
Other	2 (10)
BASELINE CHARACTERISTICS	
Time from biopsy, years	3.9 (0.9, 11.8)
Blood pressure (mmHg) – Systolic	128 (116, 132)
– Diastolic	82 (77, 86)
BMI	26.2 (24.8, 29.2)
Brain Natriuretic Peptide (pg/mL)	12.5 (8.8, 42.0)
UPCR (g/g), First morning void at screening	0.6 (0.5, 0.7)
24-hour UPCR (g/g)	0.8 (0.7, 1.1)
24-hour urine protein excretion (g/day)	1.2 (0.9, 1.5)
Urine protein excretion (g/day) ≥ 1, n (%)	14 (70)
eGFR (mL/min/1.73 m <sup>2</sup> ) <sup>‡</sup>	46 (37, 74)
Concurrent RASi, n (%)	20 (100)
ACEi	8 (40)
ARB	12 (60)

<sup>‡</sup>eGFR by CKD-EPI, Chronic Kidney Disease Epidemiology Collaboration

### Safety and Tolerability

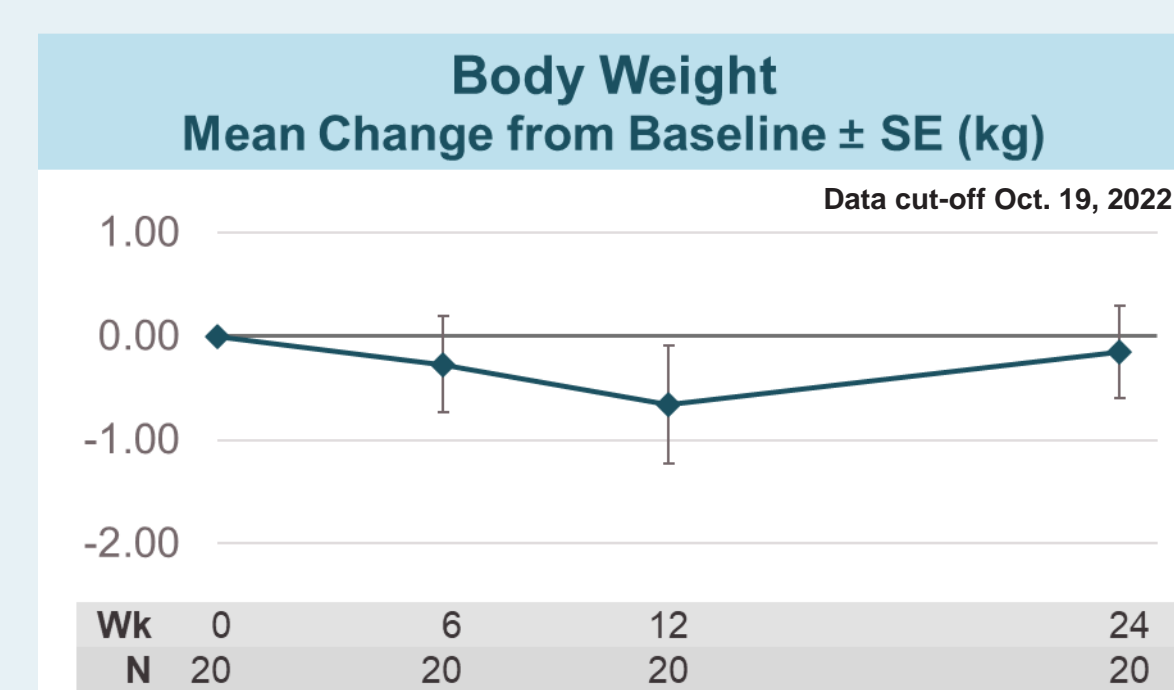
- Atrasentan was generally well-tolerated with no treatment-related severe AEs to date
- One treatment-emergent AE (headache) led to study withdrawal

AE Category (N=20)		n (%)
Treatment emergent AEs (TEAEs), Severe AEs	Subjects with any TEAE	16 (80)
	Any TEAE occurring in N>1 subjects	
	COVID-19	7 (35)
	Dizziness	3 (15)
	Peripheral edema	2 (10)
	Headache	2 (10)
	Any Moderate TEAE	6 (30)
Treatment-related AEs	Any Severe TEAE	0 (0)
	TEAE leading to discontinuation (headache)	1 (5)
	Serious AE (traffic accident unrelated to study drug)	1 (5)
	Any treatment-related AE	5 (25)
	Moderate related AEs	3 (15)
	Headache	1
Creatinine increase/Renal impairment	1	
Peripheral edema	1	

Data cut-off Oct. 19, 2022

### No Evidence of Significant Fluid Retention

- No increase in mean body weight
- No significant elevation in BNP (median change of 2.9 pg/mL at week 12)
- No meaningful change in systolic or diastolic BP
- Minimal acute change in eGFR (0.15 mL/min/1.73 m<sup>2</sup> averaged across Weeks 2 and 6)

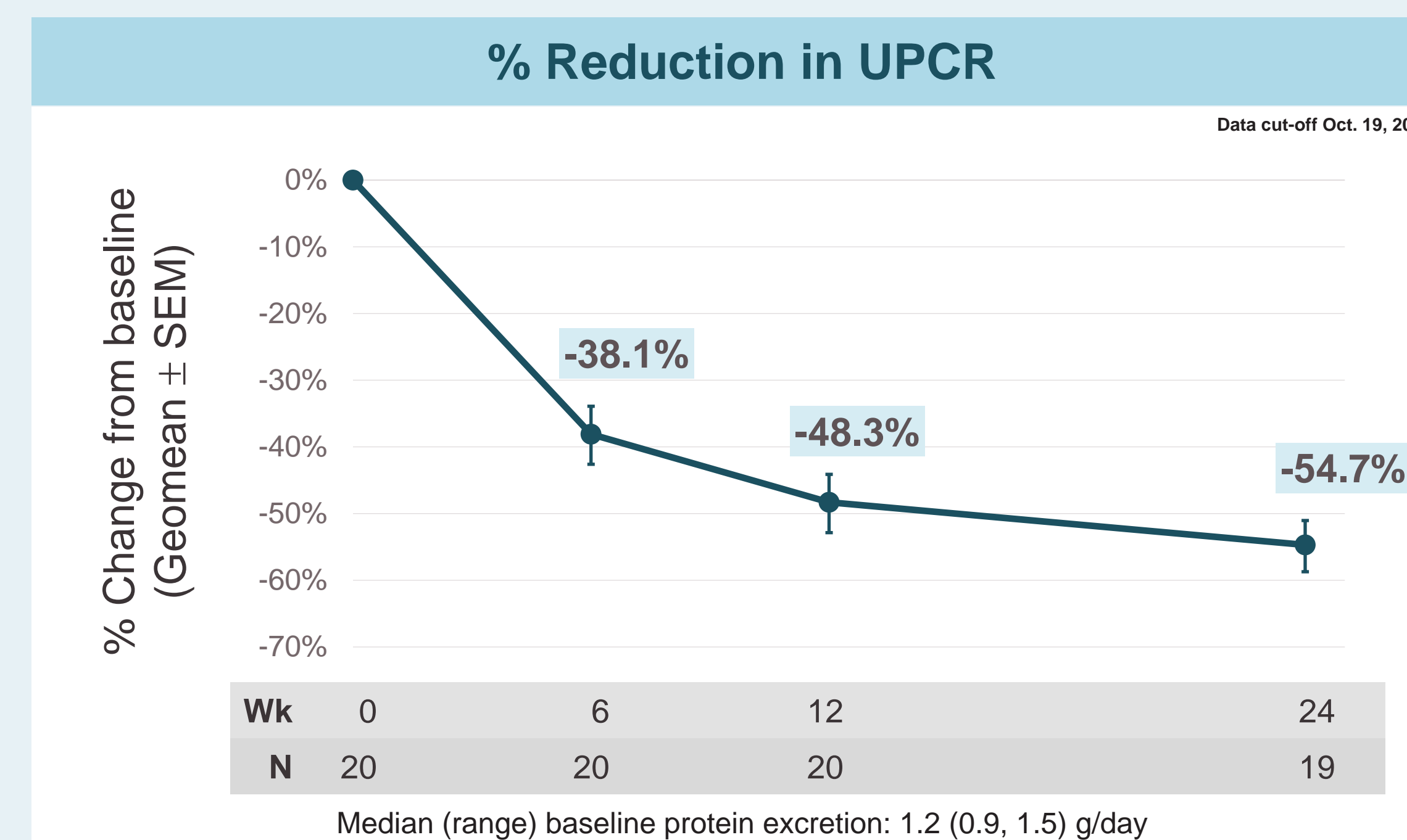


## Results

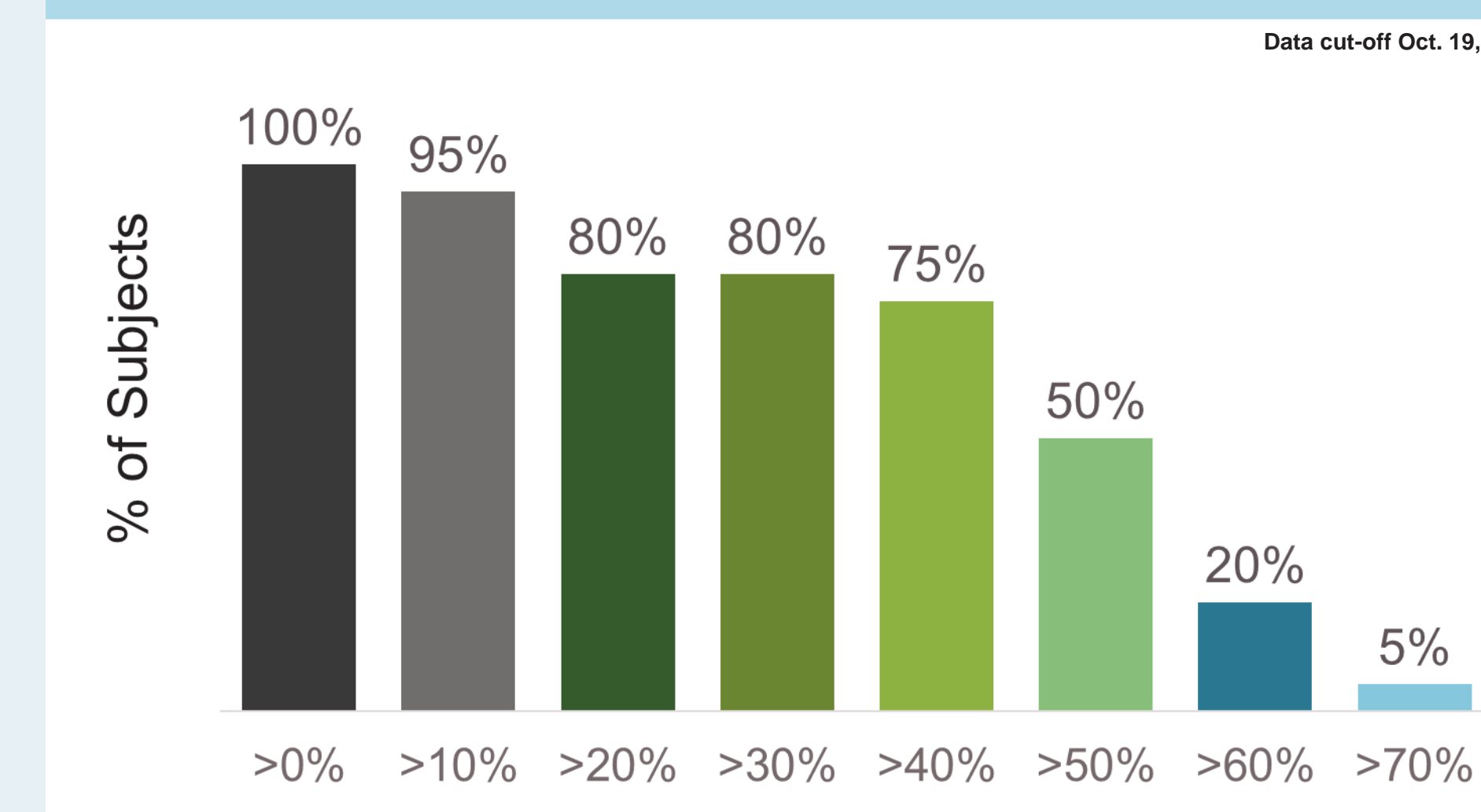
### Proteinuria Reduction in Patients with IgAN

**Treatment with atrasentan results in a durable and clinically meaningful proteinuria reduction in patients with IgAN receiving optimized standard-of-care**

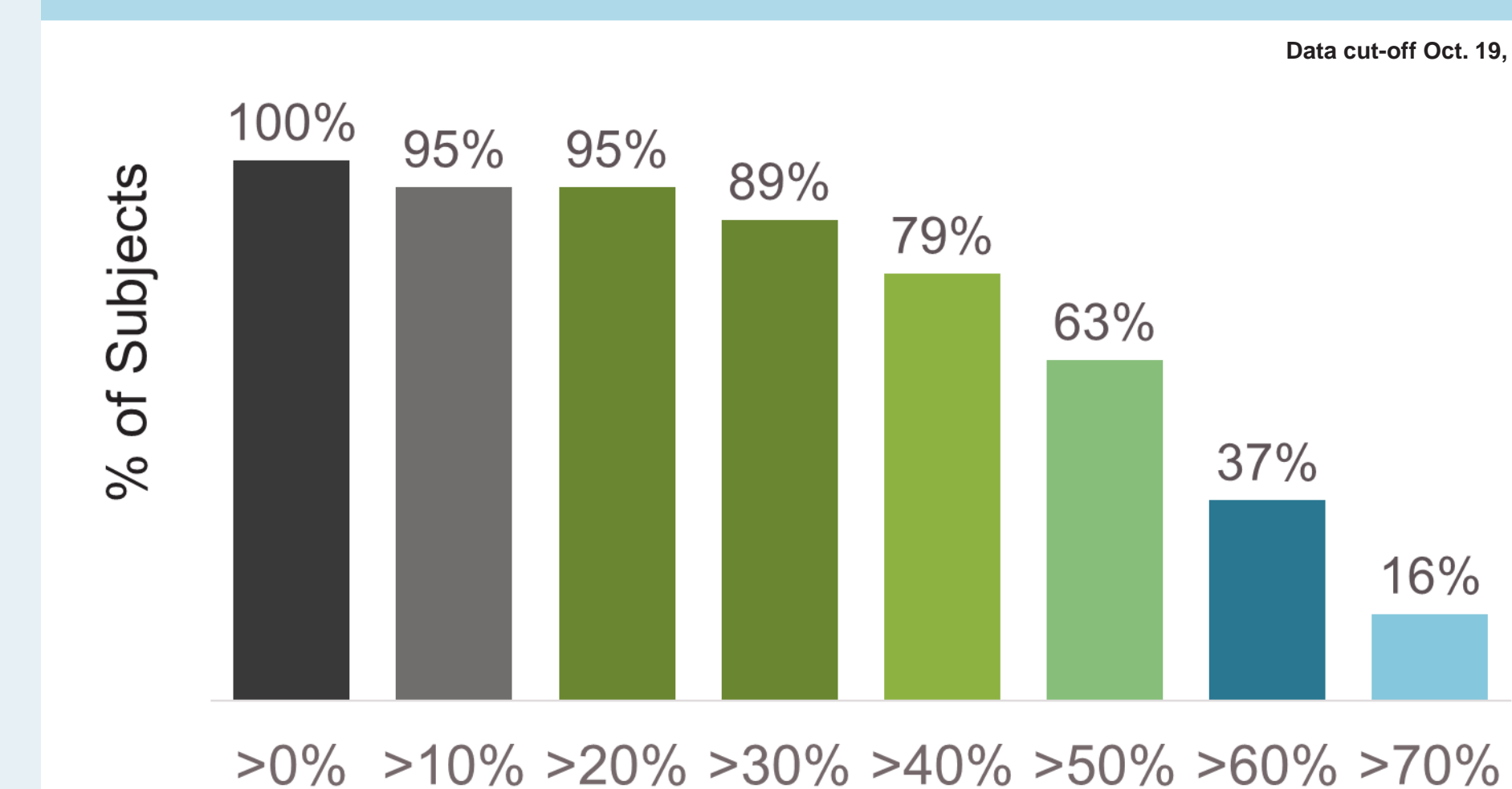
- 79% of patients achieved >40% reduction in proteinuria at Week 24



### Week 12 (n=20) UPCR Cumulative % Reduction



### Week 24 (n=19) UPCR Cumulative % Reduction



## Summary/Conclusion

- In this Phase 2 study of 20 patients with biopsy-proven IgAN, 70% of patients had baseline total urine protein >1 g/day despite optimized SOC treatment, representing an IgAN population at high risk for kidney disease progression
- Treatment with atrasentan resulted in clinically meaningful reductions in proteinuria at weeks 6, 12 and 24
- There were no meaningful changes in blood pressure nor acute eGFR changes, suggesting proteinuria reductions were not primarily due to hemodynamic effects of atrasentan
- Atrasentan was generally well-tolerated with no treatment-related SAEs
- There was no increase in BNP or mean bodyweight, suggesting minimal fluid retention

**This analysis demonstrates that treatment with atrasentan results in clinically meaningful proteinuria reductions in patients with IgAN who remain at risk for progression with residual proteinuria despite optimized standard-of-care treatment.**

### Ongoing ALIGN phase 3 trial of atrasentan in patients with IgAN

The ALIGN study (NCT04573478) is a currently enrolling/ongoing global, phase 3, randomized, double-blind, placebo-controlled study of atrasentan in patients with IgAN who are at high risk of kidney function loss.

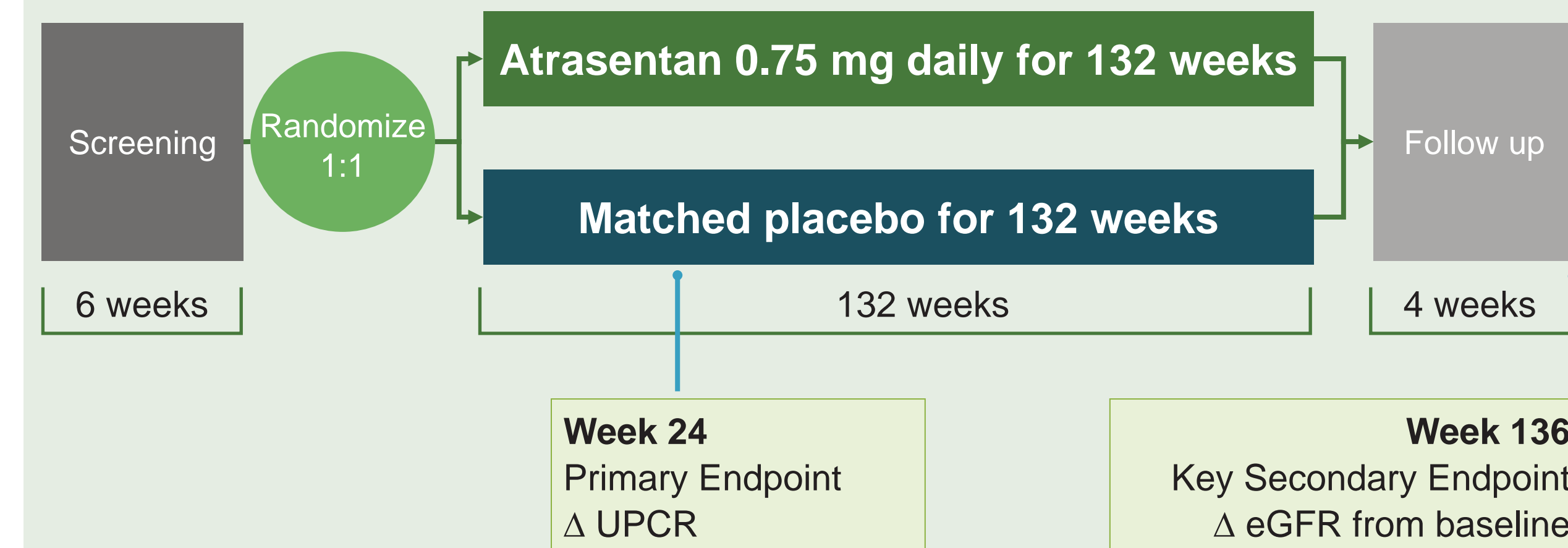
Approximately 320 patients will be enrolled across North America, South America, Europe, and Asia-Pacific.

#### Major Inclusion Criteria:

- Biopsy-proven IgAN with total protein excretion ≥ 1 g per 24 hrs and eGFR ≥ 30 mL/min/1.73 m<sup>2</sup>
- Receiving max-tolerated and optimized dose of RASi for at least 12 weeks prior to screening; a limited number of patients (up to 5%) that are unable to tolerate RASi therapy may be enrolled
- An additional stratum of up to 64 patients receiving a stable dose of SGLT2i for at least 12 weeks will be enrolled

#### Key Study Endpoints:

- Non-SGLT2i Stratum-
  - The primary endpoint is change in proteinuria from baseline at Week 24
  - The key secondary endpoint is change in eGFR from baseline at Week 136
- Additional endpoints include safety, tolerability, and quality of life



### References

- McGrogan et al., 2011, NDT; 2. Reich et al., 2007, JASN; 3. Moriyama et al., 2014, PLOS ONE; 4. Rauen et al., 2020, Kidney Int; 5. Hastings et al., 2018, Kidney Int Rep; 6. Thompson et al., 2019, CJASN; 7. Barbour et al., 2019, JAMA Int Med; 8. Inker et al., 2016, AJKD; 9. Inker et al., 2019, CJASN; 10. Tycova et al., 2018, Physiol Rev; 11. Kohan et al., 2014, Kidney Int; 12. Raina et al., 2020, Kidney Dis; 13. Lehrke et al., 2001, JASN; 14. Zanatta et al., 2012, Renal Failure; 15. Wessale et al., 2002, Clin Sci; 16. Sasser et al., JASN, 2007; 17. Olson et al., 2022, ERA; 18. Cox et al., 2021, Podocyte; 19. King et al., 2021, WCN