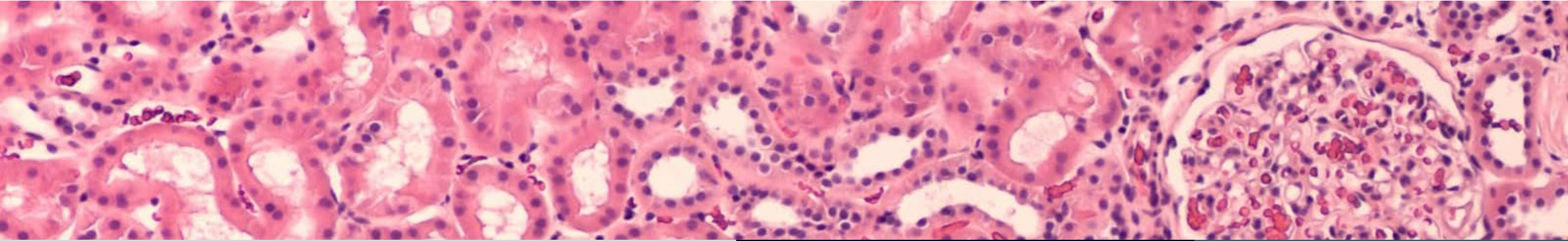


Precision's Recent Alport Experience (Rare Disease)



Natural History Study & Phase II Program

- | | |
|--|--|
| <ul style="list-style-type: none"> ➤ 7 Countries, 19 Sites ➤ 160 Patients (NH) ➤ Full-service | <ul style="list-style-type: none"> ➤ 9 Countries, 20 Sites ➤ 30 Patients (POC) ➤ Full-service |
|--|--|



Collaboration with
Advocacy Essential



Alport Experiences Shared:

- Need for visit format flexibility and incorporation into study design
 - Visits occurred in clinic and through home health services - can impact site and country selection
- Comprehensive patient recruitment strategy is critical to this disease for effective enrolment as traditional approaches are ineffective...
 - Study website recommended
 - Advocacy groups / disease foundation collaboration essential
 - Patient registration organizations
 - Outreach through social media such as Facebook is necessary
 - Involve IRB/ECs early in process so they do not become a rate limiter in recruitment strategy deployment

