Study to Assess Real-world Effectiveness of Belimumab for Treatment of Adults With LN

NCT06527872

Status	RECRUITING
Sponsor	GlaxoSmithKline
Enrollment	300 participants

Key Eligibility Criteria

Inclusion (9)

- Participants to provide a signed informed consent at the time of enrollment per protocol,
- Male or female aged 18 or over at initiation of belimumab,
- Participants received belimumab in any formulation (subcutaneous or intravenous) for the treatment of active LN prescribed as per local label in combination with standard immunosuppressive therapy/ies at initiation of belimumab,
- Participants initiated belimumab 6 to 24 months prior to study enrollment,
- Accessibility of medical records starting at belimumab initiation (including accessibility of medical records for the prior 12 months and confirmatory biopsy at any time prior to belimumab initiation),

... and 4 more (see full listing online)

Exclusion (6)

- Participants receiving renal replacement therapy (i.e., dialysis, kidney transplant, or those in end-stage kidney disease) at initiation of belimumab,
- Participant is concomitantly receiving another SLE targeted monoclonal antibody (MAb), or a MAb expected to compromise immune responses, at initiation of belimumab,
- Participants in a clinical trial during the observation period (with the exception of allowing participation in other non-interventional studies),
- Participant is pregnant at the initiation of belimumab,
- Participant with a kidney transplant at the initiation of belimumab,

... and 1 more (see full listing online)

Locations (5 total)

GSK Investigational Site, Baltimore, Maryland, United States
GSK Investigational Site, Charlotte, North Carolina, United States
GSK Investigational Site, Columbus, Ohio, United States
... and 2 more locations