

Benign Hematology Areas of Interest

(Investigator Initiated Studies and Research Collaborations)

REGN3918 (C5) pozelimab REGN3918 (C5) pozelimab + cemdisiran (ALN-CC5)	
<p>Interventional proposals in Myasthenia Gravis (MG):</p> <ul style="list-style-type: none"> • <u>Mechanistic</u> <ul style="list-style-type: none"> – Biomarkers leading to better characterization, management, and treatment options – Complement levels during disease progression • <u>Special Populations</u> <ul style="list-style-type: none"> – Triple-negative – Seronegative – Myasthenic crisis – Poor responders on non-C5 inhibiting therapies • <u>Patient Monitoring</u> <ul style="list-style-type: none"> – Geocaching – Wearable devices to monitor disease or patient Quality of Life (QoL) • <u>Indirect Treatment Comparisons/Switching Data</u> <ul style="list-style-type: none"> – Early intervention with pozelimab – Switching within the C5 inhibitor class – Switching off rituximab, IVIg, or PLEX • <u>Observational Studies</u> <ul style="list-style-type: none"> – Epidemiology – Real World Evidence (RWE) – Registries – Database 	<p>Interventional proposals in Paroxysmal Nocturnal Hemoglobinuria (PNH):</p> <ul style="list-style-type: none"> • <u>Mechanistic</u> <ul style="list-style-type: none"> – Biomarkers leading to better patient stratification – Complement levels during disease progression – Drug interactions • <u>Special Populations</u> <ul style="list-style-type: none"> – Poor responders on non-C5 inhibiting therapies – Patients that have recovered (natural resolution) – Breakthrough hemolysis • <u>Patient Monitoring</u> <ul style="list-style-type: none"> – Geocaching – Wearable devices to monitor disease or patient QoL • <u>Indirect Treatment Comparisons/Switching Data</u> <ul style="list-style-type: none"> – Switching within the C5 inhibitor class – Switching to a C5 inhibitor from non-C5 inhibiting therapies – Switching off rituximab, IVIg, or PLEX • <u>Observational Studies</u> <ul style="list-style-type: none"> – Epidemiology – RWE – Registries – Database – Patient Reported Outcomes (PRO) in patients using the PNH Specific Questionnaire
<p>Note that Regeneron cannot currently accept any proposals for Investigator Sponsored Studies or Research Collaborations using REGN3918 (C5) pozelimab and REGN3918 (C5) pozelimab + cemdisiran (ALN-CC5) in the European Union</p>	