

- S1 Real-World Effectiveness of Mepolizumab in Patients with Chronic Rhinosinusitis with Nasal Polyps: A Pre/Post Assessment of Oral Corticosteroid Use and Sinus Surgery in the US
Funded by GSK [218957]
- S2 Reduction of Systematic Corticosteroid (SCS) Related Complications with Mepolizumab in Severe Asthma: A Real-World Assessment
Funded by GSK (218950)
- S3 Real-world Impact of Treated Hereditary Angioedema Attacks on Patients' Employment and Work Productivity
Funded by KalVista Pharmaceuticals
- S4 Real-World Impact of Treated Hereditary Angioedema Attacks on Patients' Quality of Life
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- S5 Sebetralstat for On-demand Treatment of Hereditary Angioedema Attacks: US Subgroup Analysis from the Double-blind, Placebo-controlled Phase 3 KONFIDENT Trial
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- S6 Real-World Effectiveness of Berotralstat in HAE With and Without C1-Inhibitor Deficiency
Funded by BioCryst Pharmaceuticals
- S7 EPOPEX, Efficacy and Safety of Epicutaneous Immunotherapy in Peanut-Allergic Toddlers: 1-Year Open-Label Extension to EPITOPE
Funded by DBV Technologies
- S8 Remibrutinib in Chronic Spontaneous Urticaria: Efficacy and Safety at Week 24 from the REMIX-1/-2 Studies
Funded by Novartis Pharma AG, Basel, Switzerland.
- S9 Prevalence and Incidence of Diagnosed Chronic Spontaneous Urticaria (CSU) in the United States (US), Treatment Patterns and Disease Control
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- S10 Coexisting Allergic Rhinitis in Patients with Moderate-To-Severe Asthma Initiating Dupilumab In Real-World Clinical Practice: The RAPID Registry Study
Funded by Sanofi and Regeneron Pharmaceuticals Inc.
- S11 Dupilumab Improves Health-Related Quality of Life and Work Productivity Among Adults with Moderate-to-Severe Atopic Dermatitis in Clinical Practice: 4-Year Follow-up Results from the RELIEVE-AD Study
Funded by Sanofi and Regeneron Pharmaceuticals Inc.
- S12 Dupilumab Treatment Reduces Signs in Patients with Atopic Hand and Foot Dermatitis: Results from a Phase 3, Randomized, Double-Blind, Placebo-Controlled Trial
Funded by Sanofi and Regeneron Pharmaceuticals Inc.
- S13 Dupilumab Induces Clinical Remission in Children with Uncontrolled, Moderate-to-Severe, Type 2 Inflammatory Asthma
Funded by Sanofi and Regeneron Pharmaceuticals Inc.
- S14 Mepolizumab Reduced Steroid Burden for Patients With Eosinophilic Granulomatosis With Polyangiitis With And Without A Vasculitic Phenotype
Funded by GSK (115921; NCT02020889)
- S15 Real-World Mepolizumab Use in Eosinophilic Granulomatosis with Polyangiitis (EGPA) Demonstrates Improvement in End-Organ Manifestations and Long-Term Corticosteroid Benefit: An Analysis of Allergy Practices in the United States
Funded by GSK [218960]
- S16 Efficacy and safety of amltelimab (an OX40 ligand antibody) in patients with moderate-to-severe atopic dermatitis: 52-week results from a Phase 2b trial (STREAM-AD)
Funded by Kymab LTD, a Sanofi Company
- S17 Dupilumab Reduces Inflammatory Biomarkers in Patients Aged 6 Months to 18 Years with Moderate-to-Severe or Severe Atopic Dermatitis
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- S18 Dupilumab Is Efficacious in Patients With Eosinophilic Esophagitis Regardless Of Prior History Of Comorbidities
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- S19 Dupilumab normalized the expression of genes dysregulated in eosinophilic esophagitis (EoE) in esophageal biopsies from a clinical trial of children aged 1–11 years
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- S20 Effect of Benralizumab versus Mepolizumab on Reduction in Oral Glucocorticoid Use in Patients with Eosinophilic Granulomatosis with Polyangiitis: Phase 3 MANDARA Study
Funded by Astra Zeneca
- S21 Complete Remission in Eosinophilic Granulomatosis with Polyangiitis (EGPA) in the MANDARA Trial of Benralizumab versus Mepolizumab
Funded by Astraneca
- S22 Lebrikizumab Maintains Improvements in the Patient-Oriented Eczema Measure Through 2 Years of Treatment in Patients with Moderate-to-Severe Atopic Dermatitis
Funded by: Lilly

- S23 Is Asthma Clinical Remission Achievable by Inhaled Therapy? A Post Hoc Analysis of Single Inhaler Triple Therapy with FF/UMEC/VI in the CAPTAIN Trial
Funded by: GSK (205715; NCT02924688)
- S24 Relief and Resolution of Attack Symptoms Following On-Demand Treatment with a Single Dose of Oral Bradykinin B2 Receptor Antagonist Deucricitabant Immediate-Release Capsule in Patients with Hereditary Angioedema
Funded by RAPIDE-1 is a Pharvaris
- S25 13.2 mg Intranasal Epinephrine Spray Demonstrates Comparable PK/PD and Safety to 0.3 mg Epinephrine Autoinjector
Funded by Bryn Pharma
- S26 Pharmacokinetic and Pharmacodynamic Effects of 13.2 mg Intranasal Epinephrine Treatment in Congestion
Funded by Bryn Pharma
- S27 Phase 2 OLE Two-Year Analysis of Donidalorsen Taken Every 4 Weeks or Every 8 Weeks in Patients with Hereditary Angioedema
Funded by IONIS Pharmaceuticals
- S28 Updated Results of a Phase 1a Trial of STAR-0215 for Hereditary Angioedema
Funded by Astria Therapeutics
- S29 Avapritinib Led to Reductions in Symptom Burden and Polypharmacy in Patients with Indolent Systemic Mastocytosis (ISM)
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- S30 C1 Esterase Inhibitor (C1-INH) Response as a Supportive Diagnostic Criterion for Patients with Suspected Hereditary Angioedema with Normal C1-INH
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- S31 Association Between Nickel and Propolis on Patch Testing
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