Sno	REC Reference	IRAS	Name of Trial	Target Number of Patients Agreed?	Minimum Number of Patients Agreed	Maximum Number of Patients Agreed	Target Date To Recruit Patients Agreed?	Date Agreed to Recruit Target Number of Patients	Total Number of Patients Recruited at the Agreed Target Date	Date That The Trial Closed to Recruitment	Total Number Of Study Participants Recruited	Reason for Closure of Trial
1	14/ES/0001	141631	Randomised, Double-Blind, Placebo-Controlled, Parallel-Group Study to Assess Cardiovascular Outcomes Following Treatment with Ertugliflozin (MK-8835/PF-04971729) in Participants with Type 2 Diabetes Mellitus and Established Vascular Disease	Number Agreed	10	10	Not Available / Not agreed		0	29/11/2019	3	Recruitment Finished
2	14/WM/0049	145163	An openlabel singlearm multicentre noncontrolled phase 3a trial investigating safety and efficacy of nonacog betapegol (N9GP) in prophylaxis and treatment of bleeding episodes in previously untreated patients with haemophilia B (FIX activity ≤2%)	Range Agreed	1	1	Date Agreed	30/09/2019	0	02/03/2020	0	Withdrawn By Host
3	15/LO/1419	183975	A randomized, phase 3 trial with anti-PD-1 monoclonal antibody pembrolizumab (MK-3475) versus placebo for patients with early stage NSCLC after resection and completion of standard adjuvant therapy (PEARLS)	Range Agreed	5	8	Date Agreed	15/11/2019	11	15/11/2019	11	Recruitment Finished
4	15/YH/0478	186697	A Phase 3, Double-Blind, Randomized, Long-Term, Placebo-Controlled, Multicenter Study Evaluating the Safety and Efficacy of Obeticholic Acid in Subjects with Nonalcoholic Steatohepatitis	Range Agreed	1	4	Date Agreed	30/06/2019	4	17/06/2019	4	Recruitment Finished
5	16/LO/0086	193561	A Single-Masked, Randomised, Controlled, Parallel Group, Phase 3 Clinical Trial Of Retinal Gene Therapy For Choroideremia Using An Adeno-Associated Viral Vector (AAV2) Encoding Rab Escort Protein 1 (REP1)	Range Agreed	1	10	Date Agreed	30/09/2019	3	02/10/2019	3	Recruitment Finished
6	16/LO/0718	203652	A phase 3, multicentre, single-arm, open-label study of the efficacy and safety of B-domain deleted recombinant porcine factor VII (BAX802) in subjects with congenital haemophilia A with factor VIII inhibitors undergoing surgical or other invasive procedures	Range Agreed	1	2	Date Agreed	01/07/2020	0	22/05/2019	0	Withdrawn By Host

7	16/LO/2126	218039	A randomized trial comparing the ELUVIA™ drug-eluting stent versus bare metal self-expanding nitinol stents in the treatment of superficial femoral and/or proximal popliteal arteries.	Range Agreed	1	10	Date Agreed	31/01/2020	1	31/03/2020	1	Recruitment Finished
8	16/NW/0082	184887	Breast reconstruction Outcomes With and without StratticE	Range Agreed	100	200	Date Agreed	31/10/2019	121	31/10/2019	121	Recruitment Finished
9	16/SC/0551	214620	XIRIUS Study (A Dose Escalation, Phase 1/2 Clinical Trial of Retinal Gene Therapy for X-linked Retinitis Pigmentosa Using an Adeno-Associated Viral Vector (AAV8) Encoding Retinitis Pigmentosa GTPase Regulator (RPGR))	Range Agreed	1	15	Date Agreed	28/02/2020	7	20/03/2020	7	Withdrawn By Sponsor
10	16/SC/0615	210762	A Phase 4 open-label randomized controlled study COmparing the effectiveness of adalimumab iNTROduction andmethotrexate dose escaLation in subjects with Psoriatic Arthritis (CONTROL)	Range Agreed	1	5	Date Agreed	15/10/2018	11	07/05/2019	12	Recruitment Finished
11	16/WM/0511	218115	A Phase 3, Prospective, Multicenter, Uncontrolled, Open-Label Clinical Study to Determine the Efficacy, Safety, and Tolerability of rVWF with or without ADVATE in the Treatment and Control of Bleeding Episodes, the Efficacy and Safety of rVWF in Elective and Emergency Surgeries, and the Pharmacokinetics (PK) of rVWF in Children Diagnosed with Severe von Willebrand Disease.	Number Agreed	1	1	Date Agreed	31/12/2019	0	23/04/2019	0	Recruitment Finished
12	17/EE/0474	229785	An Open-Label, Randomized, Multi-Center, Parallel Group, Two-Arm Study to Assess the Safety, Overall Tolerability, and Antiviral Activity of Brincidofovir versus Standard of Care for Treatment of Adenovirus Infections in High-Risk Pediatric Allogeneic Hematopoietic Cell Transplant Recipients.	Range Agreed	1	4	Date Agreed	31/05/2019	3	09/05/2019	3	Withdrawn By Sponsor
13	17/EM/0063	213979	A Phase 3 Multicenter, Randomized, Double-Blind, Placebo-Controlled Trial of the FLT3 Inhibitor Gilteritinib (ASP2215) Administered as Maintenance Therapy Following Induction/Consolidation Therapy for Subjects with FLT3/ITD AML in First Complete Remission	Range Agreed	1	3	Date Agreed	31/03/2019	0	10/04/2019	0	Withdrawn By Sponsor

14	17/EM/0122	224376	A Phase 3, Open-Label, Multicenter Study Of ALXN1210 In Children And Adolescents With Atypical Hemolytic-Uremic Syndrome (aHUS)	Range Agreed	1	1	Date Agreed	30/11/2020	0	23/01/2020	0	Recruitment Finished
15	17/EM/0152	216307	A Multi-center, Randomised, Double-blind, Placebo-controlled Phase III Trial of the FLT3 Inhibitor Gilteritinib Administered as Maintenance Therapy Following Allogeneic Transplant for Patients with FLT3/ITD AML	Range Agreed	1	3	Date Agreed	31/03/2020	5	15/02/2020	5	Recruitment Finished
16	17/EM/0404	223875	A Phase 3, Multi-center, Randomized, Double-Masked Study to Evaluate the Clinical Efficacy and Safety of SHP640 (PVP-Iodine 0.6% and Dexamethasone 0.1%) Ophthalmic Suspension Compared to PVP-Iodine and Placebo in the Treatment of Adenoviral Conjunctivitis	Range Agreed	1	10	Date Agreed	31/12/2020	2	03/05/2019	2	Withdrawn By Sponsor
17	17/LO/0041	219676	CardioMEMS Heart Failure System "Outside US" Post-Approval Study	Range Agreed	1	10	Date Agreed	01/09/2019	7	21/06/2019	7	Recruitment Finished
18	17/LO/0236	220765	PROMINENT PEMAFIBRATE TO REDUCE CARDIOVASCULAR OUTCOMES BY REDUCING TRIGLYCERIDES IN PATIENTS WITH DIABETES	Range Agreed	5	10	Date Agreed	31/03/2020	6	14/02/2020	6	Recruitment Finished
19	17/LO/0243	219613	A randomized, double-blind, multi-dose, placebo-controlled study to evaluate the efficacy, safety and tolerability ofGSK2330672 administration for the treatment of pruritus in patients with primary biliary cholangitis.(GLIMMER:GSK2330672 triaL of Ibat inhibition with Multidose Measurement for Evaluation of Response).	Range Agreed	1	4	Date Agreed	24/09/2019	0	24/09/2019	0	Recruitment Finished
20	17/LO/0372	220433	A phase III randomized open-label multi-center study of ruxolitinib vs. best available therapy in patients with corticosteroid-refractory acute graft vs. host disease after allogenic stem cell transplantation	Range Agreed	1	3	Date Agreed	07/05/2019	3	07/05/2019	3	Recruitment Finished
21	17/LO/1088	228921	A Phase 3, 2-Arm, Open-label Study to Evaluate the Safety and Pharmacodynamics of Long-termIvacaftor Treatment in Subjects With Cystic Fibrosis Who Are Less Than 24 Months of Age at Treatment Initiation and Have a CFTR Gating Mutation	Range Agreed	1	3	Date Agreed	11/12/2019	0	11/12/2019	0	Recruitment Finished

22	17/LO/1306	228268	A phase III randomized open-label multi-center study of ruxolitinib vs. best available therapy in patients with corticosteroid-refractory chronic graft vs host disease after allogenic stem cell transplantation (REACH 3)	Range Agreed	1	3	Date Agreed	11/10/2019	4	06/12/2019	4	Recruitment Finished
23	17/LO/1930	235819	A randomized, controlled, open-label, multiple ascending dose study of intravenous brincidofovir in adult allogeneic hematopoietic cell transplant recipients with adenovirus viremia	Range Agreed	1	2	Date Agreed	30/09/2019	0	09/05/2019	0	Withdrawn By Sponsor
24	17/NE/0200	220486	Strategic MAnagement to Optimize Response To CardiacResynchronization Therapy Registry (SMART Registry)	Number Agreed	12	12	Date Agreed	31/08/2019	27	19/08/2019	27	Recruitment Finished
25	17/NE/0358	220871	Relative bioavailability and comparative pharmacokinetics of 13-CRA oral liquid and extracted capsule formulations: a randomised, open label, multi-dose, cross-over clinical trial in patients requiring treatment cycles of 13-CRA.	Range Agreed	2	3	Date Agreed	30/11/2019	2	17/07/2019	2	Recruitment Finished
26	17/NI/0096	225743	HOPE-1 (HYDRATION FOR OPTIMAL PULMONARY EFFECTIVENESS)	Range Agreed	2	4	Date Agreed	10/11/2018	2	01/04/2019	3	Withdrawn By Host
27	17/NS/0106	230907	The hydrus microstent for refractory open-angle glaucoma: A prospective, multicenter clinical trial	Range Agreed	3	10	Date Agreed	31/05/2019	3	23/10/2019	3	Recruitment Finished
28	17/NW/0209	213231	An open-label ascending dose cohort study to assess the safety, pharmacokinetics, and preliminary efficacy ofneoGAA (GZ402666) in patients with infantile-onset Pompe disease treated with alglucosidase alfa who demonstrateclinical decline or sub-optimal clinical response	Range Agreed	1	3	Date Agreed	19/04/2019	1	15/04/2019	1	Recruitment Finished
29	17/NW/0247	222303	A Phase 3, Multicenter, Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Efficacy	Number Agreed	5	5	Date Agreed	31/03/2025	1	30/09/2019	1	Recruitment Finished
30	17/NW/0399	224760	Study of the measurement of volatile agents to diagnose urological disease	Range Agreed	90	150	Date Agreed	31/07/2019	82	31/07/2019	82	Recruitment Finished

31	17/SC/0462	228889	A Randomized, Double-blind Phase 1/2a Study to Evaluate the Safety, Tolerability and Immunogenicity of Ad26.RSV.preF in Adults 18 to 50 Years of Age and RSV-Seropositive Toddlers 12 to 24 Months of Age	Number Agreed	4	4	Date Agreed	31/03/2018	0	14/06/2019	4	Recruitment Finished
32	17/SC/0554	225522	A Phase III, multicenter, randomized controlled study to compare safety and efficacy of a haploidentical HSCT and adjunctive treatment with ATIR101, a T-lymphocyte enriched leukocyte preparation depleted ex vivo of host alloreactive T-cells, versus a haploidentical HSCT with post-transplant cyclophosphamide in patients with a hematologic malignancy (HATCY study)	Number Agreed	4	4	Date Agreed	30/12/2020	4	12/11/2019	4	Withdrawn By Sponsor
33	17/SW/0221	232448	A randomized, partially-blinded, active-controlled,multicenter study of secukinumab to demonstrate reduction of radiographic progression versus GP2017 (adalimumab biosimilar) at 104 weeks and to assess the long term safety, tolerability and efficacy up to 2 years in patients with active ankylosing spondylitis	Number Agreed	4	4	Date Agreed	03/11/2021	0	16/05/2019	0	Withdrawn By Sponsor
34	17/WA/0347	234208	A randomised, double-blind, placebo-controlled, parallel-group, multicentre study to demonstrate the effects ofSotagliflozin on Cardiovascular and Renal Events in Patients with Type 2 Diabetes, Cardiovascular Risk Factors andModerately Impaired Renal Function	Number Agreed	12	12	Date Agreed	31/12/2019	12	16/09/2019	12	Recruitment Finished
35	17/WM/0308	226910	Randomized, Double-Blind, Phase 3B Trial to Evaluate the Safety and Efficacy of 2 Treatment Regimens of Aztreonam 75 mg Powder and Solvent for Nebulizer Solution / Aztreonam for Inhalation Solution (AZLI) in Pediatric Subjects with Cystic Fibrosis (CF) and New Onset Respiratory Tract Pseudomonas aeruginosa (PA) Infection/Colonization	Range Agreed	1	2	Date Agreed	01/11/2019	2	17/02/2020	2	Recruitment Finished

36	17/WM/0316	226628	A Post-Approval Registry of the TREO® Stent-Graft for Patients with Infrarenal Abdominal Aortic Aneurysms	Range Agreed	1	10	Date Agreed	01/06/2020	4	31/12/2019	4	Recruitment Finished
37	17/YH/0426	231118	A Randomized, Double-Blind, Placebo-Controlled Study to Investigate the Efficacy and Safety of Dupilumab Administered Concomitantly with Topical Corticosteroids in Patients, ≥6 Years to <12 Years Of Age, with Severe Atopic Dermatitis	Range Agreed	2	20	Date Agreed	30/09/2018	5	21/08/2019	5	Recruitment Finished
38	17/YH/0432	236091	A double blind (sponsor open) placebo-controlled, stratified, parallelgroup study to evaluate the efficacy and safety of repeat doses of GSK3772847 inparticipants with moderate to severe asthma with allergic fungal airway disease (AFAD).	Range Agreed	3	5	Date Agreed	28/06/2019	0	30/06/2019	0	Withdrawn By Sponsor
39	18/EE/0005	232671	A Multicenter, Open-Label Study To Estimate The Effect Sizes Of HRCT Endpoints In Response To Glucocorticoid Induction Therapy In Subject With Pulmonary Sarcoidsis	Number Agreed	2	2	Date Agreed	31/08/2019	0	04/09/2019	0	Withdrawn By Sponsor
40	18/EM/0153	240773	A PHASE 3, RANDOMIZED, DOUBLE-BLIND,PLACEBO-CONTROLLED STUDY TO EVALUATE THEEFFICACY AND SAFETY OF AG-348 IN NOTREGULARLY TRANSFUSED ADULT SUBJECTS WITHPYRUVATE KINASE DEFICIENCY	Range Agreed	1	2	Date Agreed	28/02/2020	0	05/02/2020	0	Recruitment Finished
41	18/EM/0365	252018	An open-label extension trial of the long-term safety of nintedanib in patients with Progressive Fibrosing Interstitial Lung Disease (PF-ILD	Number Agreed	3	3	Date Agreed	31/07/2019	3	31/07/2019	3	Recruitment Finished
42	18/LO/0190	227705	Clinical investigation of the eyeWatch glaucoma drainage device	Number Agreed	15	15	Date Agreed	31/05/2019	15	31/05/2019	15	Recruitment Finished
43	18/LO/0656	240062	Multiple escalating dose study of BAY 1093884 in adults with Haemophilia A or B with or without inhibitors	Number Agreed	1	1	Date Agreed	16/07/2020	1	05/08/2019	1	Withdrawn By Sponsor
44	18/LO/0711	241788	A Phase 3, Randomized, Double-Blind, Placebo-Controlled, 12-month Study to Evaluate the Efficacy and Safety of MK-7264 in Adult Participants with Chronic Cough (PN030)	Number Agreed	8	8	Date Agreed	07/08/2019	5	12/08/2019	5	Recruitment Finished

45	18/LO/0712	241782	A Phase 3, Randomized, Double-Blind, Placebo-Controlled, 12-Month Study to Evaluate the Efficacy and Safety of MK-7264 in Adult Participants with Chronic Cough (PN027)	Number Agreed	8	8	Date Agreed	07/08/2019	8	17/05/2019	8	Recruitment Finished
46	18/LO/0755	245151	A Randomized, Open-label, 24-Week Safety, Efficacy, and Pharmacokinetic Study of Teduglutide in Infants 4 to 12 Months of Age with Short Bowel Syndrome Who are Dependent on Parenteral Support	Range Agreed	1	2	Date Agreed	30/11/2019	1	02/03/2020	2	Recruitment Finished
47	18/LO/1007	242697	Patient-Reported Outcomes with the Accu-Chek® SoloMicropump System vs. Insulet OmniPod® vs. Multiple DailyInjection Therapy in Type 1 Diabetes	Range Agreed	1	4	Date Agreed	31/07/2019	4	31/07/2019	4	Recruitment Finished
48	18/LO/1311	248599	A phase III, multicenter, randomized, double-masked, active comparatorcontrolled study to evaluate the efficacy and safety of RO6867461 in patients with diabetic macular edema (RHINE)	Number Agreed	2	2	Date Agreed	31/12/2019	2	22/08/2019	2	Recruitment Finished
49	18/LO/1922	251251	A Phase 2a, Open-label, Single and Multiple Dose Study to Evaluate the Pharmacokinetics, Safety, Tolerability and Treatment Effect of GBT440 in Pediatric Participants with Sickle Cell Disease	Range Agreed	1	3	Date Agreed	31/05/2021	2	06/01/2020	2	Withdrawn By Sponsor
50	18/NE/0102	240702	A phase I, open–label, randomized, pharmacokinetic, pharmacodynamic, and safety study ofEtrolizumab followed by open–label extension and safety monitoring in paediatric patientsFrom 4 years to less than 18 years of age with moderate to severe ulcerative colitis orModerate to severe crohn's disease	Number Agreed	1	1	Date Agreed	31/12/2019	2	20/08/2019	2	Recruitment Finished
51	18/NE/0142	242919	A Phase 3, Double-Blind, Randomized, Placebo-Controlled, Multicenter Study to Evaluate the Efficacy and Safety of Obeticholic Acid in Subjects with Compensated Cirrhosis due to Nonalcoholic Steatohepatitis Protocol 747-304	Range Agreed	1	8	Date Agreed	08/11/2019	1	22/11/2019	1	Recruitment Finished

52	18/NE/0169	243977	A 6-month randomised, double-blind, placebo controlled multicentre parallel group study to evaluate efficacy and safety of bumetanide 0.5mg twice a day followed by an open label active 6-month treatment period with bumetanide (0.5mg twice a day) and a 6 weeks discontinuation period after treatment stop	Range Agreed	6	12	Date Agreed	31/03/2020	11	21/02/2020	11	Recruitment Finished
53	18/NE/0221	249183	A Phase I, Multicenter, Open-Label, Single-Dose, Dose-Ranging Study to Assess the Safety and Tolerability of SB-913, a rAAV2/6-based Gene Transfer in Subjects with Mucopolysaccharidosis II (MPSII)	Range Agreed	1	4	Date Agreed	31/08/2019	0	25/04/2019	0	Withdrawn By Sponsor
54	18/NI/0178	253727	A Phase 1 / 2, Drug-Drug Interaction Study of FDL169 and FDL176 in Healthy Subjects and in Cystic Fibrosis Subjects Homozygous for the F508del-CFTR mutation	Range Agreed	1	4	Date Agreed	29/02/2020	0	10/12/2019	0	Withdrawn By Sponsor
55	18/NI/0240	258468	An Observational, Prospective Multicentre Clinical Study to assess the safety and clinical performance of a New Single-use Negative Pressure Wound Therapy System (PICO 7Y) for the Simultaneous Management of Bilateral Closed Incisions in Oncoplastic Breast Surgery Patients	Number Agreed	10	10	Date Agreed	31/12/2019	13	18/05/2019	13	Recruitment Finished
56	18/NW/0098	242717	Two-part, double-blind, placebo-controlled, randomized, parallel-group study: (Part 1) in healthy male subjects to assess safety and tolerability of ascending repeated oral doses of BAY 1902607 including its effect on the pharmacokinetics of a sub-therapeutic dose of midazolam (MDZ), followed by (Part 2) a two-way crossover administration of four different doses of BAY 1902607 in patients with refractory chronic cough to assess safety, tolerability and efficacy for proof of concept	Number Agreed	2	2	Date Agreed	30/04/2019	2	05/07/2019	3	Recruitment Finished
57	18/NW/0215	239446	A Phase 2, Randomized, Placebo-Controlled Study Of Safety And Efficacy Following Repeat-Dose Administration Of Evinacumab (Anti-Angptl3) In Patients With Severe Hypertriglyceridemia (Shtg) At Risk For Acute Pancreatitis	Range Agreed	1	3	Date Agreed	30/11/2019	4	09/07/2019	4	Recruitment Finished

58	18/NW/0325	244842	A Randomized, Double-Blind, Placebo-Controlled Study of the Safety and Efficacy of Varying Doses and Dose Regimens of Evinacumab in Patients with Persistent Hypercholesterolemia Despite Maximally Tolerated Lipid Modifying Therapy	Range Agreed	1	3	Date Agreed	31/12/2019	3	24/11/2019	3	Recruitment Finished
59	18/NW/0565	246158	A Phase II randomised , double blind, placebo-controlled, parallel group, multicentre study to evaluate the safety and efficacy of repeated oral doses of Blautix in adult subjects with irritable bowel syndrome (IBS) subtypes IBS-C and IBS-D	Range Agreed	6	14	Date Agreed	31/03/2020	4	17/01/2020	4	Withdrawn By Host
60	18/NW/0599	242748	Study of TRE Seals on Early Post-operative Subjects (STEPS)	Range Agreed	1	10	Date Agreed	29/07/2019	3	30/10/2019	4	Recruitment Finished
61	18/NW/0742	236247	A Phase 2, Randomized, Double-Blind, Placebo-Controlled Efficacy and Safety Study of Palovarotene in Subjects with Multiple Osteochondromas	Range Agreed	4	15	Date Agreed	30/06/2019	3	05/12/2019	7	Withdrawn By Host
62	18/SC/0395	245499	A Multicenter, Randomized, Phase III Registration Trial of Transplantation of NiCord®, Ex Vivo Expanded, Umbilical Cord Blood-derived, Stem and Progenitor Cells, versus Unmanipulated Umbilical Cord Blood for Patients with Hematological Malignancies	Range Agreed	1	3	Date Agreed	31/12/2019	1	24/12/2019	1	Recruitment Finished
63	18/SS/0140	252391	Epidemiological analysis for Hereditary Angioedema Disease (EHA Study)	Number Agreed	100	100	Date Agreed	30/06/2022	11	28/02/2020	11	Withdrawn By Sponsor
64	18/SW/0162	248769	A prospective follow up study to assess performance, safety and efficacy of the PICO 7 NPWT system for surgically closed incision sites and skin grafts	Range Agreed	5	8	Date Agreed	29/03/2019	9	21/05/2019	9	Recruitment Finished
65	18/WM/0022	237623	AN OPEN-LABEL STUDY TO EVALUATE THEEFFICACY AND SAFETY OF AG-348 IN REGULARLYTRANSFUSED ADULT SUBJECTS WITH PYRUVATE KINASE (PK) DEFICIENCY	Range Agreed	1	2	Date Agreed	31/12/2019	1	31/12/2019	1	Recruitment Finished
66	18/WM/0204	244427	A Phase 3, Open-label Study Evaluating the Long-term Safety and Efficacy of VX-445 Combination Therapy in Subjects With Cystic Fibrosis Who Are Homozygous or Heterozygous for the F508del Mutation	Range Agreed	1	3	Date Agreed	31/08/2019	3	18/04/2019	3	Recruitment Finished

67	18/WM/0312	228906	ACCEPTABILITY STUDY ON KETOGENIC DIET SUPPLEMENT USING CAMBROOKE KETOVIE™ MEDICAL FOODS	Range Agreed	1	15	Date Agreed	30/09/2019	11	30/09/2019	11	Recruitment Finished
68	18/YH/0055	241341	A Phase 2, Prospective, Randomized, Open-label Study on the Efficacy of Defibrotide Added to Standard of Care Immunoprophylaxis for the Prevention of Acute Graft-versus- Host-Disease in Adult and Pediatric Patients After Allogeneic Hematopoietic Stem Cell Transplant	Range Agreed	1	2	Date Agreed	31/12/2019	4	31/10/2019	4	Recruitment Finished
69	18/YH/0136	214921	A multi-centre, pilot, prospective, trial of DermaRep™ Device in the treatment of venous leg ulcers	Range Agreed	1	5	Date Agreed	30/09/2019	6	30/09/2019	6	Recruitment Finished
70	19/EM/0003	251874	A Randomized, Double-Blind, Placebo Controlled, Global Phase 3 Study Of Edasalonexent In Pediatric Patients With Duchenne Muscular Dystrophy	Range Agreed	1	3	Date Agreed	30/09/2019	5	30/09/2019	5	Recruitment Finished
71	19/EM/0047	257111	A Phase III, Multicenter, Randomized, Double-Masked, Active Comparator-Controlled Study To Evaluate The Efficacy And Safety Of Faricimab In Patients With Neovascular Age-Related Macular Degeneration (TENAYA)	Number Agreed	4	4	Date Agreed	31/12/2019	3	24/10/2019	3	Recruitment Finished
72	19/EM/0094	259038	Effects of N-Acetyl-L-Leucine on Niemann-Pick disease type C: A multinational, multicenter, open-label, rater-blinded Phase II study.	Range Agreed	1	4	Date Agreed	31/01/2020	1	31/01/2020	1	Recruitment Finished
73	19/LO/0157	256597	A study to evaluate safety, reactogenicity and immunogenicity of GSK Biologicals' RSV investigational vaccine based on viral proteins encoded by chimpanzee-derived adenovector (ChAd155-RSV) (GSK3389245A) in infants	Number Agreed	2	2	Date Agreed	30/09/2019	1	20/10/2019	1	Recruitment Finished
74	19/LO/0351	256743	A Phase 3, Randomized, Double-Blind, Study Comparing Risankizumab to Placebo in Subjects with Active Psoriatic Arthritis (PsA) Who Have a History of Inadequate Response to or Intolerance to at Least One Disease Modifying Anti- Rheumatic Drug (DMARD) Therapy	Range Agreed	1	3	Date Agreed	30/06/2020	0	11/03/2020	0	Recruitment Finished

75	19/LO/0352	257478	A Phase 3, Randomized, Double-Blind Study Comparing Risankizumab to Placebo in Subjects with Active Psoriatic Arthritis Including Those Who Have a History of Inadequate Response or intolerance to Biologic Therapy(ies)	Range Agreed	1	3	Date Agreed	30/06/2020	0	28/11/2019	0	Withdrawn By Sponsor
76	19/NW/0026	249432	VX18-121-101: A Phase 2, Randomized, Double-blind, Controlled Study to Evaluate the Safety and Efficacy of VX-121 Combination Therapy in Subjects Aged 18 Years and Older with Cystic Fibrosis	Range Agreed	1	4	Date Agreed	30/09/2019	6	30/08/2019	6	Recruitment Finished
77	19/NW/0482	269475	A Phase 3b, Randomized, Double-blind, Controlled Study Evaluating the Efficacy and Safety of VX-445/Tezacaftor/Ivacaftor in Cystic Fibrosis Subjects, Homozygous for F508del	Range Agreed	1	5	Date Agreed	30/03/2020	5	19/12/2019	5	Withdrawn By Sponsor
78	19/SC/0031	257722	A Randomized, Placebo-controlled, Double-blind, Phase 1/2a Study to Evaluate the Safety, Reactogenicity, and Immunogenicity of Ad26.RSV.preF in RSV-seronegative Toddlers 12 to 24 Months of Age	Range Agreed	1	5	Date Agreed	30/11/2019	0	06/08/2019	0	Recruitment Finished