

| REC Ref No. | IRAS No. | Name of Trial  | Target Number of Participants Available? | Recruitment Target | Target Date To Recruit Participants Available? | Date Agreed To Recruit Target Number of Participants | Total Number of Participants Recruited at The Agreed Target Date | Date Study Closed to Recruitment | Recruitment Total | RTT Met? | Recruitment Closure Reason |
|-------------|----------|--|--|--------------------|--|--|--|----------------------------------|-------------------|----------|----------------------------|
| 20/WM/0123  | 282099   | A randomized, double-blind, placebo-controlled, multi-centre study to evaluate the safety and efficacy of tocilizumab in patients with severe COVID-19 pneumonia   | Number Agreed                            | 3                  | Date Agreed                                    | 29/01/2021   | 3  | 28/05/2020                       | 12                | Yes      | Recruitment Finished       |
| 20/NE/0105  | 282026   | A Phase 3 Randomized Study to Evaluate the Safety and Antiviral Activity of Remdesivir (GS-5734, Ñϕ) in Participants with Moderate COVID-19 Compared to Standard of Care Treatment   | Number Agreed                            | 4                  | Date Agreed                                    | 04/06/2020   | 3  | 29/05/2020                       | 3                 | Yes      | Recruitment Finished       |
| 20/NE/0104  | 282007   | A Phase 3 Randomized Study to Evaluate the Safety and Antiviral Activity of Remdesivir (GS-5734, Ñϕ) in Participants with Severe COVID-19  | Number Agreed                            | 9                  | Date Agreed                                    | 29/05/2020   | 6  | 29/05/2020                       | 6                 | Yes      | Recruitment Finished       |
| 20/YH/0090  | 278137   | AN OPEN-LABEL, MULTICENTER, ROLLOVER STUDY TO EVALUATE THE SAFETY AND TOLERABILITY OF LONG-TERM ADMINISTRATION OF GANTENERUMAB IN PARTICIPANTS WITH ALZHEIMER, Åôs DISEASE   | Number Agreed                            | 2                  | Date Agreed                                    | 07/09/2020   | 2  | 30/09/2020                       | 2                 | Yes      | Recruitment Finished       |
| 18/NS/0145  | 254786   | A NON-INTERVENTIONAL, MULTICENTER, MULTIPLE COHORT STUDY INVESTIGATING THE OUTCOMES AND SAFETY OF ATEZOLIZUMAB UNDER REAL-WORLD CONDITIONS IN PATIENTS TREATED IN ROUTINE CLINICAL PRACTICE  | Number Agreed                            | 2                  | Date Agreed                                    | 31/12/2020   | 2  | 31/12/2020                       | 2                 | Yes      | Recruitment Finished       |
| 19/LO/1566  | 267041   | A Randomized, Placebo-controlled, Phase 2 Study to Evaluate the Safety and Pharmacodynamics of Once-daily Oral IW-1701 in Patients with Stable Sickle Cell Disease   | Number Agreed                            | 1                  | Date Agreed                                    | 30/09/2020   | 0  | 06/04/2020                       | 0                 | No       | Recruitment Finished       |
| 19/EE/0164  | 261589   | T-cell Lymphoma anti-KIR3DL2 therapy. An open label, multi-cohort, multi-center phase II study evaluating the efficacy and safety of IPH4102 alone or in combination with chemotherapy in patients with advanced T-cell lymphoma.  | Number Agreed                            | 3                  | Date Agreed                                    | 07/09/2020   | 0  | 12/05/2020                       | 0                 | No       | Withdrawn By Sponsor       |
| 19/LO/1263  | 263485   | A Randomized, Double-Blind, Controlled Phase 3 Study of Cabozantinib in Combination with Nivolumab and Ipilimumab versus Nivolumab and Ipilimumab in Subjects with Previously Untreated Advanced or Metastatic Renal Cell Carcinoma of Intermediate or Poor Risk   | Number Agreed                            | 3                  | Date Agreed                                    | 01/12/2020   | 1  | 01/12/2020                       | 1                 | No       | Recruitment Finished       |
| 19/EE/0201  | 264655   | A Randomised, Single-blind, Placebo-controlled, Phase 1b Single Ascending (Part A) and Multiple Dose (Part B) First-in-Man Study in Adult Patients with Non-transfusion-Dependent Beta-Thalassaemia or Low Risk Myelodysplastic Syndrome to Investigate the Safety, Tolerability, Pharmacokinetic and Pharmacodynamic Response of SLN124 | Number Agreed                            | 3                  | Date Agreed                                    | 03/05/2021   | 0  | 20/03/2020                       | 0                 | No       | Withdrawn By Sponsor       |

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| 18/NE/0360 | 254062 | An open-label, non-randomised study on efficacy, pharmacokinetics, pharmacodynamics, safety and tolerability of LNP023 in two patient populations with C3 glomerulopathy   | Number Agreed | 2  | Date Agreed | 30/06/2020 | 1 | 30/06/2020 | 1 | No  | Recruitment Finished |
| 19/NW/0174 | 261666 | A Phase III Randomized, Double-Blind, Placebo-Controlled, Multi-Regional, International Study of Durvalumab in Combination with Gemcitabine plus Cisplatin versus Placebo in Combination with Gemcitabine plus Cisplatin for Patients with First-Line Advanced Biliary Tract Cancers                     | Number Agreed | 3  | Date Agreed | 31/12/2020 | 6 | 31/12/2020 | 6 | Yes | Recruitment Finished |
| 19/EM/0020 | 252705 | A phase II, open-label, prospective, single-arm, study to assess ability of eltrombopag to induce sustained remission in subjects with ITP who are refractory or relapsed after first-line steroids (TAPER)  | Number Agreed | 2  | Date Agreed | 30/11/2020 | 3 | 30/11/2020 | 3 | Yes | Recruitment Finished |
| 19/LO/0150 | 258058 | A double-blind, placebo-controlled, randomised phase III trial to assess the safety and efficacy of Viaskin Peanut in peanut-allergic young children 1-3 years of age  | Number Agreed | 10 | Date Agreed | 30/07/2020 | 7 | 31/07/2020 | 7 | No  | Recruitment Finished |
| 17/LO/1921 | 236323 | A PHASE 3, RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDY TO ASSESS THE EFFICACY AND SAFETY OF BIVV009 IN PATIENTS WITH PRIMARY COLD AGGLUTININ DISEASE WITHOUT A RECENT HISTORY OF BLOOD TRANSFUSION  | Number Agreed | 1  | Date Agreed | 31/03/2020 | 1 | 31/03/2020 | 1 | Yes | Recruitment Finished |
| 18/EE/0099 | 239606 | An open label, randomised, phase III Study comparing trifluridine/tipiracil (S 95005) in combination with bevacizumab to capecitabine in combination with bevacizumab in first-line treatment of patients with metastatic colorectal cancer who are not candidate for intensive therapy (SOLSTICE study) | Number Agreed | 10 | Date Agreed | 31/03/2020 | 2 | 19/08/2020 | 2 | No  | Recruitment Finished |
| 19/YH/0013 | 255993 | A Randomized, Controlled Phase 3 Study of Cabozantinib (XL184) in Combination with Atezolizumab versus Sorafenib in Subjects with Advanced Hepatocellular Carcinoma Who have not received Previous Systemic Anticancer therapy.  | Number Agreed | 3  | Date Agreed | 01/04/2023 | 3 | 19/08/2020 | 3 | Yes | Recruitment Finished |
| 19/LO/1183 | 259840 | IPX203-B16-02  | Number Agreed | 3  | Date Agreed | 30/07/2020 | 1 | 30/07/2020 | 3 | Yes | Recruitment Finished |
| 18/LO/1258 | 248312 | Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Efficacy and Safety of BG00011 in Patients With Idiopathic Pulmonary Fibrosis   | Number Agreed | 2  | Date Agreed | 30/09/2020 | 1 | 31/12/2020 | 1 | No  | Withdrawn By Sponsor |
| 19/SC/0350 | 266934 | A Phase 1B, Randomized, Subject- and Investigator-Blinded, Placebo-Controlled, Multi-Center Clinical Trial to Evaluate the Safety and Pharmacokinetics of Inhaled GB002, and Assess Changes in Imaging and Biomarkers in Subjects with WHO Group 1 Pulmonary Arterial Hypertension (PAH)                 | Number Agreed | 2  | Date Agreed | 31/10/2020 | 0 | 31/10/2020 | 0 | No  | Recruitment Finished |
| 18/LO/1558 | 248946 | A Phase 2 Trial of Pembrolizumab (MK-3475) in Combination with Platinum Doublet Chemotherapy and Radiotherapy for Participants with Unresectable, Locally Advanced Stage III Non-Small Cell Lung Cancer (NSCLC)  | Number Agreed | 2  | Date Agreed | 27/02/2020 | 0 | 27/02/2020 | 0 | No  | Withdrawn By Sponsor |
| 17/LO/0232 | 220795 | An Open-Label, Randomized, Crossover Trial utilizing a Single-Blinded Rater to evaluate APL-130277 compared to s.c. Apomorphine in Levodopa Responsive Subjects with Parkinson's Disease Complicated by Motor Fluctuations   | Number Agreed | 4  | Date Agreed | 30/04/2020 | 1 | 30/04/2020 | 2 | No  | Recruitment Finished |

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| 18/WA/0276 | 246112 | Randomised Double-Blind, Placebo-Controlled (within dose groups) and Active Controlled (Eplerenone group) Trial to Investigate the Safety, Tolerability, Pharmacokinetics and Pharmacodynamics of 4 Oral Doses of BI 690517 Over 28 Days in Female and Male Patients with Diabetic Nephropathy | Number Agreed | 2  | Date Agreed | 30/10/2020 | 0  | 01/10/2020 | 0  | No  | Recruitment Finished |
| 18/NW/0765 | 250220 | Validating a Device for the Detection of Bacteria and Leukocytes in Peritoneal Effluent from Patients with Suspected Peritonitis   | Number Agreed | 10 | Date Agreed | 02/11/2020 | 13 | 31/12/2020 | 13 | Yes | Recruitment Finished |
| 18/SW/0219 | 250410 | A PHASE 2, OPEN-LABEL, MULTICENTER STUDY TO DETERMINE THE EFFICACY, SAFETY, PHARMACOKINETICS, AND PHARMACODYNAMICS OF AG-348 IN ADULT SUBJECTS WITH NON-TRANSFUSION-DEPENDENT THALASSEMIA-AG348-C-010  | Number Agreed | 1  | Date Agreed | 31/01/2020 | 1  | 10/03/2020 | 1  | Yes | Recruitment Finished |
| 16/LO/1567 | 242092 | A Randomized, Controlled, Open-label, Global Phase 3 Study Comparing the Efficacy of the anti-PD-1 Antibody BGB-A317 versus Chemotherapy as Second Line Treatment in Patients with Advanced Unresectable/Metastatic Esophageal Squamous Cell Carcinoma   | Number Agreed | 3  | Date Agreed | 31/08/2020 | 5  | 09/04/2020 | 5  | Yes | Recruitment Finished |
| 17/LO/0182 | 213821 | A Prospective, Randomized, Multicenter Controlled Trial of CERAMENT, Nϕ G as Part of Surgical Repair of Open Diaphyseal Tibial Fractures   | Number Agreed | 5  | Date Agreed | 31/03/2020 | 5  | 15/06/2020 | 4  | Yes | Recruitment Finished |
| 18/ES/0071 | 241904 | A Multi-Center, Randomized, Assessor-Blind, Controlled Trial Comparing the Occurrence of Major Adverse Cardiovascular Events in Patients with Prostate Cancer and Cardiovascular Disease Receiving Degarelix or Leuprolide   | Number Agreed | 5  | Date Agreed | 29/10/2020 | 2  | 24/03/2020 | 1  | No  | Recruitment Finished |
| 18/LO/1215 | 245798 | A Randomized, Phase 3 Study of Eryaspase in Combination with Chemotherapy versus Chemotherapy Alone as Second-Line Treatment in Patients with Pancreatic Adenocarcinoma<br>TRYbeCA-1 ,Äi TRial of erYaspase in pancreatic CANcer   | Number Agreed | 2  | Date Agreed | 30/08/2020 | 2  | 30/08/2020 | 2  | Yes | Recruitment Finished |
| 18/LO/1882 | 253030 | A Phase 2, Randomized, Double-Blind, Placebo-Controlled Study of Abiraterone Acetate plus Prednisone with or without Abemaciclib in Patients with Metastatic Castration-Resistant Prostate Cancer  | Number Agreed | 2  | Date Agreed | 16/11/2020 | 2  | 23/10/2020 | 2  | Yes | Recruitment Finished |
| 18/NE/0214 | 245998 | MN39158 - A Long Term Extension Study in Multiple Sclerosis  | Number Agreed | 3  | Date Agreed | 01/06/2020 | 3  | 01/06/2020 | 3  | Yes | Recruitment Finished |
| 18/NI/0129 | 218623 | Efficacy and safety assessment of T4032 (unpreserved bimatoprost 0.01%) versus Lumigan-Æ 0.01% in ocular hypertensive or glaucomatous patients.  | Number Agreed | 4  | Date Agreed | 01/05/2020 | 1  | 01/05/2020 | 1  | No  | Withdrawn By Sponsor |
| 18/LO/1007 | 242697 | Patient-Reported Outcomes with the Accu-Chek√Ç-Æ Solo Micropump System vs. Multiple Daily Injection Therapy vs. mylife OmniPod√Ç-Æ in Patients with Type 1 Diabetes  | Number Agreed | 4  | Date Agreed | 31/05/2020 | 0  | 29/05/2020 | 0  | No  | Recruitment Finished |
| 18/NW/0514 | 249725 | SELECT - Semaglutide effects on cardiovascular outcomes in people with overweight or obesity   | Number Agreed | 35 | Date Agreed | 23/10/2020 | 32 | 23/10/2020 | 32 | No  | Recruitment Finished |

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| 18/LO/1187 | 240011 | (A Multicenter, Randomized, Double-Blind, Placebo-Controlled Phase 3 Study in Ovarian Cancer Patients Evaluating Rucaparib and Nivolumab as Maintenance Treatment following Response to Front-Line Platinum-Based Chemotherapy)   | Number Agreed | 6 | Date Agreed | 23/09/2023 | 4 | 30/01/2020 | 4 | No  | Recruitment Finished |
| 18/LO/0430 | 241907 | An Adaptive, Open-Label, Dose-Finding, Phase 1/2 Study Investigating the Safety, Pharmacokinetics, and Clinical Activity of PRN1008, an Oral BTK Inhibitor, in Patients with Relapsed Immune Thrombocytopenia   | Number Agreed | 1 | Date Agreed | 31/12/2020 | 3 | 31/12/2020 | 3 | Yes | Recruitment Finished |
| 18/YH/0099 | 241430 | A randomized, double-blind, placebo-controlled, parallel group study to evaluate the efficacy and safety of CNP520 in participants at risk for the onset of clinical symptoms of Alzheimer,Ås Disease (AD)  | Number Agreed | 5 | Date Agreed | 26/03/2020 | 4 | 26/03/2020 | 4 | No  | Withdrawn By Sponsor |
| 18/SC/0392 | 244109 | Phase 2 Multicenter, Double-Blind, Dose Finding, Placebo Controlled, Safety, Efficacy and Pharmacokinetic Study of CXA-10 on Stable Background Therapy in Subjects with Pulmonary Arterial Hypertension (PAH)   | Number Agreed | 2 | Date Agreed | 31/08/2020 | 1 | 02/07/2020 | 1 | No  | Withdrawn By Sponsor |
| 18/SS/0010 | 238051 | A Phase 3 Study of Erdafitinib Compared With Vinflunine or Docetaxel or Pembrolizumab in Subjects with Advanced Urothelial Cancer and Selected FGFR Gene Aberrations  | Number Agreed | 1 | Date Agreed | 27/11/2020 | 1 | 27/11/2020 | 1 | Yes | Recruitment Finished |
| 18/LO/0235 | 240315 | An Open Label Extension Study of GBT440 Administered Orally to Patients with Sickle Cell Disease Who Have Participated in GBT440 Clinical Trials  | Number Agreed | 1 | Date Agreed | 01/03/2020 | 3 | 01/03/2020 | 3 | Yes | Recruitment Finished |
| 18/EM/0153 | 240773 | A Phase 3, Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Efficacy and Safety of AG-348 in Not Regularly Transfused Adult Subjects With Pyruvate Kinase Deficiency  | Number Agreed | 1 | Date Agreed | 17/09/2020 | 2 | 28/02/2020 | 2 | Yes | Recruitment Finished |
| 18/LO/0070 | 238482 | A Pivotal Clinical Trial of the Management of th Medically-Refractory Dyskinesia Symptoms or Motor Fluctuations of Advanced Idiopathic Parkinson,Ås Disease With Unilateral Lesioning of the Globus Pallidum Using the ExAblate Neuro System  | Number Agreed | 6 | Date Agreed | 02/03/2020 | 0 | 27/02/2020 | 0 | No  | Recruitment Finished |
| 18/WM/0022 | 237623 | AG-348 in Regularly Transfused Adult Subjects with PK Deficiency  | Number Agreed | 1 | Date Agreed | 27/09/2020 | 1 | 27/09/2020 | 1 | Yes | Recruitment Finished |
| 17/EE/0474 | 229785 | The open-label, randomised, multi-centre, 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 paediatric allogeneic haematopoietic cell transplant recipients | Number Agreed | 5 | Date Agreed | 21/02/2020 | 1 | 09/05/2020 | 1 | No  | Recruitment Finished |
| 17/NW/0351 | 224954 | A phase II single arm clinical trial of a Tailored ImmunoTherapy Approach with Nivolumab in subjects with metastatic or advanced Renal Cell Carcinoma   | Number Agreed | 2 | Date Agreed | 31/12/2020 | 3 | 31/12/2020 | 3 | Yes | Recruitment Finished |
| 17/LO/1149 | 220763 | A Phase 3, multicenter, randomized, double-blind, double-dummy, active-controlled study to assess the efficacy and safety of maribavir compared to valganciclovir for the treatment of cytomegalovirus (CMV) infection in hematopoietic stem cell transplant recipients.                                | Number Agreed | 2 | Date Agreed | 30/08/2021 | 2 | 06/01/2020 | 2 | Yes | Recruitment Finished |

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| 17/WM/0146  | 220303 | A Placebo-Controlled, Double-Blind, Parallel-Group, 24-Month Study to Evaluate the Efficacy and Safety of E2609 in Subjects with Early Alzheimer's Disease  | Number Agreed | 4  | Date Agreed | 30/06/2020 | 4  | 26/03/2020 | 4  | Yes | Withdrawn By Sponsor |
| 16/EM/0193  | 190690 | A phase III, double-blind, randomized placebo-controlled study to evaluate the effects of dalcetrapib on cardiovascular (CV) risk in a genetically defined population with a recent Acute Coronary Syndrome (ACS): The dal-GenE trial | Number Agreed | 6  | Date Agreed | 31/12/2020 | 6  | 31/12/2020 | 6  | Yes | Recruitment Finished |
| 04/MRE07/35 | 31586  | Systemic Therapy in Advancing or Metastatic Prostate Cancer: Evaluation of Drug Efficacy  | Number Agreed | 69 | Date Agreed | 31/12/2020 | 76 | 31/12/2020 | 76 | Yes | Recruitment Finished |