

| 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/HRA/2068 | 283394 | A Phase 3 Open-label, Randomized, Controlled Study to Evaluate the Efficacy and Safety of Intravenously Administered Ravulizumab Compared with Best Supportive Care in Patients with COVID-19 Severe Pneumonia, Acute Lung Injury, or Acute Respiratory Distress Syndrome | Number Agreed | 5 | Date Agreed | 29/06/2020 | 0 | 29/06/2020 | 0 | No | Recruitment Finished |
| 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 | Range Agreed | 3 | Date Agreed | 29/01/2021 | 12 | 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) 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) in Participants with Severe COVID-19 | Number Agreed | 6 | 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'S 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 |
| 18/WS/0187 | 242800 | Retrospective and prospective international EKoSoNic registry Of the treatment and Clinical OUTcomes of patients with Pulmonary Embolism | Number Agreed | 3 | Date Agreed | 31/03/2020 | 2 | 30/09/2020 | 2 | 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/NW/0328 | 263312 | A Phase 2 Randomized, Double-Blind, Placebo Controlled Study of the Safety and Efficacy of BMS-986165 in Subjects with Moderate to Severe Ulcerative Colitis | Number Agreed | 2 | Date Agreed | 31/12/2020 | 0 | 15/03/2021 | 0 | No | Recruitment Finished |

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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 | 3 | Yes | 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 | 30/10/2020 | 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 | 30/07/2020 | 7 | No | Recruitment Finished |
| 19/NW/0342 | 264974 | A Phase 3/4, Multinational, Double-Blind, Randomized, Placebo-Controlled Study of MGL-3196 in Patients With Non-Alcoholic Steatohepatitis (NASH) | Range Agreed | 2 | Date Agreed | 20/12/2020 | 0 | 20/12/2020 | 0 | No | 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 candidates 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 |
| 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 | 1 | 31/12/2020 | 1 | No | Recruitment Finished |
| 18/EM/0256 | 248467 | Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Multicenter, Phase 3 Study to Evaluate the Efficacy and Safety of Intravenous BIIB093 (Glibenclamide) for Severe Cerebral Edema following Large Hemispheric Infarction | Number Agreed | 3 | Date Agreed | 29/06/2020 | 0 | 30/11/2020 | 0 | No | Recruitment Finished |
| 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 | 17/03/2021 | 2 | No | Recruitment Finished |
| 18/SC/0240 | 230920 | A Phase III, Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Efficacy, and Safety Study of Gantenerumab in Patients with Early (Prodromal to Mild) Alzheimer's Disease | Number Agreed | 8 | Date Agreed | 11/01/2021 | 9 | 11/01/2021 | 9 | Yes | 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 | 29/07/2020 | 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 | 02/11/2020 | 13 | 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/WM/0117 | 223480 | Setmelanotide (RM-493) Phase 2 Treatment Trial in Patients with rare genetic disorders of obesity | Number Agreed | 3 | Date Agreed | 31/01/2021 | 2 | 31/01/2021 | 3 | Yes | Recruitment Finished |
| 17/LO/0182 | 213821 | A Prospective, Randomized, Multicenter Controlled Trial of CERAMENTâ,ç 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/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 TRYbeCA-1 â€œ TRIal of erYaspase in pancreatic CAncer | Number Agreed | 2 | Date Agreed | 30/08/2020 | 2 | 01/12/2020 | 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 LumiganÂ® 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-ChekÂ,Â® Solo Micropump System vs. Multiple Daily Injection Therapy vs. mylife OmniPodÂ,Â® in Patients with Type 1 Diabetes | Number Agreed | 4 | Date Agreed | 31/05/2020 | 0 | 31/05/2020 | 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 | 34 | 18/05/2020 | 34 | Yes | 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/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/NE/0136 | 240494 | A Phase 3, Randomized, Double-blind Study of Adjuvant Nivolumab versus Placebo for Participants with Hepatocellular Carcinoma Who Are at High Risk of Recurrence after Curative Hepatic Resection or Ablation | Number Agreed | 1 | Date Agreed | 01/05/2020 | 0 | 01/05/2020 | 0 | No | Recruitment Finished |

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| 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/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/WS/0180 | 225790 | A Phase 111b, Randomised, Double-blind, Placebocontrolled, MulticentreStudy of Olaparib Maintenance Retreatment in Patients with EpithelialOvarian Cancer Previously Treated With a PARPi and Responding toRepeat Platinum Chemotherapy (OReO) | Number Agreed | 4 | Date Agreed | 30/04/2021 | 1 | 15/01/2021 | 1 | No | Recruitment Finished |
| 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 Alzheimerâ€™s Disease | Number Agreed | 4 | Date Agreed | 30/06/2020 | 4 | 30/06/2020 | 4 | Yes | Recruitment Finished |
| 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 |