

| Research Ethics Committee Reference Number | Integrated Research Application System Number | Name of Trial | Target Number Of Patients Agreed? | Minimum Number Of Patients Agreed (Enter Same In Both If Only One Number) | Maximum Number Of Patients Agreed (Enter Same In Both If Only One Number) | 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 | Reason For Closure Of Trial |
|--|---|--|-----------------------------------|---|---|---|--|--|---|-----------------------------|
| 14/WM/0170 | 154982 | A Phase 3, Randomized Study to Evaluate the Efficacy of Momelotinib versus Best Available Therapy in Anemic or Thrombocytopenic Subjects with Primary Myelofibrosis, Post-polycythemia Vera Myelofibrosis, or Post-essential Thrombocythemia Myelofibrosis | Number Agreed | 2 | 2 | Not Available / Not Agreed | | 3 | 12/01/2016 | Recruitment Finished |
| 14/EE/1235 | 159181 | A PHASE 2 RANDOMIZED, DOUBLE BLIND PLACEBO CONTROLLED TRIAL OF MHAA4549A, A MONOCLONAL ANTIBODY, IN COMBINATION WITH OSELTAMIVIR VERSUS OSELTAMIVIR FOR TREATMENT OF SEVERE? INFLUENZA A INFECTION | Number Agreed | 2 | 2 | Not Available / Not Agreed | | 0 | 02/02/2016 | Withdrawn By Host |
| 14/NW/0119 | 148925 | A 4 week phase 2a, multicentre, randomised, double-blind, placebo-controlled add-on study into safety, tolerability and efficacy of 200 mg t.i.d. of PL37 in patients with peripheral neuropathic pain of diabetic origin treated with pregabalin or gabap | Number Agreed | 10 | 10 | Not Available / Not Agreed | | 0 | 16/03/2016 | Withdrawn By Sponsor |
| 14/SC/0033 | 146569 | PHASE 3 MULTI-CENTER, DOUBLE-BLIND, RANDOMIZED, PLACEBO-CONTROLLED, PARALLEL GROUP EVALUATION OF THE EFFICACY, SAFETY, AND TOLERABILITY OF PF-04950615, IN REDUCING THE OCCURRENCE OF MAJOR CARDIOVASCULAR EVENTS IN HIGH RISK SUBJECTS | Number Agreed | 6 | 6 | Not Available / Not Agreed | | 5 | 26/01/2016 | Recruitment Finished |
| 15/SW/0197 | 181752 | GO29432: A PHASE III, OPEN-LABEL, RANDOMIZED STUDY OF MPDL3280A (ANTI-PDL1 ANTIBODY) COMPARED WITH GEMCITABINE+CISPLATIN OR CARBOPLATIN FOR PD-L1-SELECTED, CHEMOTHERAPY NAIVE PATIENTS WITH STAGE IV SQUAMOUS NON-SMALL CELL LUNG CANCER | Number Agreed | 4 | 4 | Not Available / Not Agreed | | 0 | 19/04/2016 | Recruitment Finished |
| 14/LO/0324 | 146711 | A Multicenter, Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Safety and Efficacy of Pulmaquin? in the Management of Chronic Lung Infections with Pseudomonas aeruginosa in Subjects with Non-Cystic Fibrosis Bronchiectasis, includin | Number Agreed | 3 | 3 | Not Available / Not Agreed | | 1 | 06/01/2016 | Recruitment Finished |
| 15/LO/0967 | 178036 | iNOVATE Study: A randomized, double-blind, placebo-controlled, phase 3 study of ibrutinib or placebo in combination with Rituximab in Subjects with Waldenstr?m?s Macroglobulinemia | Number Agreed | 5 | 5 | Not Available / Not Agreed | | 0 | 13/01/2016 | Recruitment Finished |
| 14/NE/1214 | 162696 | A Randomised, OpenLabel, Multicenter, Phase 2 Trial Comparing Veliparib Plus Carboplatin and Paclitaxel Versus Investigator's Choice of Standard Chemotherapy in Subjects Receiving First Cytotoxic Chemotherapy for Metastatic or Advanced NonSquamous No | Number Agreed | 6 | 6 | Not Available / Not Agreed | | 0 | 10/06/2016 | Recruitment Finished |
| 13/SC/0490 | 135889 | A PHASE III, Randomized, double-blind, placebo-controlled study to assess the efficacy, safety, and tolerability of lebrikizumab in adolescent patients with severe uncontrolled asthma who are on inhaled corticosteroids and a second controller medicat | Number Agreed | 3 | 3 | Not Available / Not Agreed | | 2 | 30/06/2016 | Withdrawn By Sponsor |
| 13/NI/0148 | 102251 | A phase II, randomised, double blind, placebo controlled, multicentre trial to assess the oral corticosteroidsparing effect of lebrikizumab in patients with with severe corticosteroiddependant asthma | Number Agreed | 12 | 12 | Not Available / Not Agreed | | 10 | 30/06/2016 | Withdrawn By Sponsor |
| 16/LO/0026 | 195795 | A Phase 3, Randomized, Open-Label Study to Evaluate the Safety and Efficacy of Switching from Regimens Consisting of Boosted Atazanavir or Darunavir plus either Emtricitabine/Tenofovir or Abacavir/Lamivudine to GS 9883/Emtricitabine/Tenofovir Alafena | Number Agreed | 6 | 6 | Not Available / Not Agreed | | 3 | 23/06/2016 | Recruitment Finished |
| 16/LO/0029 | 195696 | A Phase 3, Randomized, Double-Blind Study to Evaluate the Safety and Efficacy of GS-9883/Emtricitabine/Tenofovir Alafenamide Versus Dolutegravir + Emtricitabine/Tenofovir Alafenamide in HIV-1 Infected, Antiretroviral Treatment-Na?ve Adults | Number Agreed | 3 | 3 | Not Available / Not Agreed | | 5 | 15/07/2016 | Recruitment Finished |

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|------------|--------|---|---------------|----|----|----------------------------|--|----|------------|----------------------|
| 15/LO/1163 | 181430 | A Phase 3b, Randomized, Double-Blind, Switch Study to Evaluate F/TAF in HIV-1 Infected Subjects who are Virologically Suppressed on Regimens Containing ABC/ETC | Number Agreed | 6 | 6 | Not Available / Not Agreed | | 6 | 29/05/2016 | Recruitment Finished |
| 13/EM/0349 | 133741 | A randomized Phase II study of Fulvestrant in combination with the dual mTOR inhibitor AZD2014 or Everolimus or Fulvestrant alone in Estrogen receptor-positive advanced or metastatic breast cancer | Number Agreed | 3 | 3 | Not Available / Not Agreed | | 1 | 13/09/2016 | Recruitment Finished |
| 13/EM0239 | 116638 | Remifentanyl Intravenous Patient Controlled Analgesia (PCA) versus Intramuscular Pethidine for Pain Relief in Labour: A Randomised Controlled Trial | Number Agreed | 40 | 40 | Not Available / Not Agreed | | 2 | 05/09/2016 | Recruitment Finished |
| 14/YH/1199 | 153953 | GALACTIC: GA101 (obinutuzumab) monoclonal Antibody as Consolidation Therapy In CLL | Number Agreed | 83 | 83 | Not Available / Not Agreed | | 4 | 23/11/2016 | Recruitment Finished |
| 14/SC/0033 | 146569 | PHASE 3 MULTI CENTER, DOUBLE BLIND, RANDOMIZED, PLACEBO CONTROLLED, PARALLEL GROUP EVALUATION OF THE EFFICACY, SAFETY, AND TOLERABILITY OF PF 04950615, IN REDUCING THE OCCURRENCE OF MAJOR CARDIOVASCULAR EVENTS IN HIGH RISK SUBJECTS | Number Agreed | 6 | 6 | Not Available / Not Agreed | | 0 | 04/11/2016 | Withdrawn By Sponsor |
| 14/SC/0032 | 142458 | PHASE 3 MULTI CENTER, DOUBLE BLIND, RANDOMIZED, PLACEBO CONTROLLED, PARALLEL GROUP EVALUATION OF THE EFFICACY, SAFETY, AND TOLERABILITY OF PF 04950615 IN REDUCING THE OCCURRENCE OF MAJOR CARDIOVASCULAR EVENTS IN HIGH RISK SUBJECTS | Number Agreed | 6 | 6 | Not Available / Not Agreed | | 0 | 04/11/2016 | Withdrawn By Sponsor |
| 14/LO/0324 | 146711 | A Multicenter, Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Safety and Efficacy of Pulmaquinr in the Management of Chronic Lung Infections with Pseudomonas aeruginosa in Subjects with Non-Cystic Fibrosis Bronchiectasis, including | Number Agreed | 3 | 3 | Not Available / Not Agreed | | | 06/01/2016 | Recruitment Finished |
| 11/LO/1570 | 80305 | CHEMO-T - Cyclophosphamide, doxorubicin, vincristine and prednisolone (CHOP) versus Gemcitabine and Cisplatin (GEM-P) in the first line treatment Of T-cell Lymphoma, a multicentre randomised phase II study | Number Agreed | 4 | 4 | Not Available / Not Agreed | | 0 | 01/12/2016 | Withdrawn By Sponsor |
| 12/LO/1078 | 97486 | A phase II randomised trial of carfilzomib, cyclophosphamide and dexamethasone (CCD) vs cyclophosphamide, velcade and dexamethasone (CVD) for first relapse or primary refractory multiple myeloma. | Number Agreed | 5 | 5 | Not Available / Not Agreed | | 8 | 06/12/2016 | Recruitment Finished |
| 13/LO/0108 | 112000 | LEGEND - A Randomised Phase II Study Comparing Lenalidomide plus Rituximab, Gemcitabine and Methylprednisolone (L-RGEM) to Rituximab, Gemcitabine, Methylprednisolone and Cisplatin (RGEMP) in Second Line Treatment of Diffuse Large B-Cell Lymphoma (DLB) | Number Agreed | 5 | 5 | Not Available / Not Agreed | | 2 | 16/12/2016 | Withdrawn By Sponsor |
| 12/NS/0093 | 112403 | Vault or Uterine prolapse surgery Evaluation. Two parallel randomised controlled trials of surgical options for upper compartment (vault or uterine) pelvic organ prolapse | Number Agreed | 10 | 10 | Not Available / Not Agreed | | 7 | 06/12/2016 | Recruitment Finished |
| 12/SC/0592 | 98570 | An Observational Post authorisation Safety Specialist Cohort Event Monitoring Study (SCEM) to Monitor the Safety and Utilization of Rivaroxaban (Xarelto) for the Prevention of Stroke in Patients with AF, Treatment of DVT and PE, and the Prevention of | Number Agreed | 30 | 30 | Not Available / Not Agreed | | 32 | 31/01/2016 | Recruitment Finished |