

**NIHR Performance Metrics - Delivering Clinical Research
Quarter 4 (2015 - 2016)**

| Research Ethics Committee Reference Number | Integrated Research Application | Name of Trial | Target Number Of Patients Agreed ? | Minimum Number Of | Maximum Number Of | Target Date To Recruit | Date Agreed to recruit target number of patients | Total Number Of Patients Recruited At | Date That The Trial Closed To Recruitment | Reason For Closure Of Trial |
|--|---------------------------------|--|------------------------------------|-------------------|-------------------|----------------------------|--|---------------------------------------|---|-----------------------------|
| 11/YH/0140 | 69800 | Luminous: Study to observe the effectiveness and safety of LUCENTIS(R) through individualised patient treatment and associated outcomes | Range Agreed | 50 | 100 | Not Available / Not Agreed | | 85 | 20/05/2015 | Recruitment Finished |
| 11/LO/1711 | 86867 | An observational study of Avastin? (Bevacizumab) in combination with chemotherapy for treatment of first line metastatic colorectal adenocarcinoma. | Not Available / Not Agreed | | | Not Available / Not Agreed | | | 02/04/2015 | Recruitment Finished |
| 12/WS/0300 | 114081 | A Randomized, DoubleBlind, PlaceboControlled, ParallelGroup Study to Evaluate the Effect of SAR236553/REGN727 on the Occurrence of Cardiovascular Events in Patients Who Have Recently Experienced an Acute Coronary Syndrome-ODYSSEY OUTCOMES | Number Agreed | 25 | 25 | Not Available / Not Agreed | | 61 | 11/11/2015 | Recruitment Finished |
| 13/SC/0268 | 124775 | An open-label, crossover, interventional Phase IV study to compare the ease of use of tobramycin inhalation powder with tobramycin inhalation solution and nebulized colistimethate for the treatment of pulmonary Pseudomonas aeruginosa in patients with cysti | Number Agreed | 5 | 5 | Not Available / Not Agreed | | 5 | 20/10/2015 | Withdrawn By Sponsor |
| 13/YH/0282 | 133239 | Openlabel, Phase IIIb study to evaluate the efficacy and safety of subcutaneous (SC) Tocilizumab monotherapy or combination therapy with methotrexate (MTX) or other nonbiologic disease modifying antirheumatic drugs (DMARDs) in patients with severe Rheumat | Number Agreed | 5 | 5 | Not Available / Not Agreed | | 7 | 05/06/2015 | Recruitment Finished |
| 13/NW/0698 | 133706 | A phase III multicenter, randomized study of oral LDK378 versus standard chemotherapy in previously untreated adult patients with ALK rearranged (ALK positive), stage IIIB or IV, non squamous nonsmall cell lung cancer (CLDK378A2301) | Number Agreed | 2 | 2 | Not Available / Not Agreed | | 7 | 15/04/2015 | Recruitment Finished |
| 13/LO/1245 | 1318223 | A Phase 1b, Multicenter, Pilot, Randomized, Double-blind Trial to Determine the Pharmacokinetics and Pharmacodynamics of Orally Administered Tolvaptan 3.75, 7.5, and 15 mg Tablets in Subjects with Syndrome of Inappropriate Antidiuretic Hormone Secretion (O | Number Agreed | 3 | 3 | Not Available / Not Agreed | | 5 | 15/06/2015 | Recruitment Finished |
| 13/NW/0612 | 135524 | A multicentre, randomised, doubleblind, parallel group, placebocontrolled, phase III efficacy and safety study of benralizumab (MEDI563) added to highdose inhaled corticosteroid plus longacting B2agonist in patients with uncontrolled asthma (SIROCCO) | Number Agreed | 1 | 1 | Not Available / Not Agreed | | 3 | 20/05/2015 | Recruitment Finished |

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| 13/YH/0389 | 142190 | A 52 week, phase 3 double-blind, randomized, placebocontrolled, parallel-group study to assess the efficacy, safety and tolerability of pf-04950615 in subjects with heterozygous familial hypercholesterolemia (B1481021) | Number Agreed | 6 | 6 | Not Available / Not Agreed | 6 | 02/02/2015 | Recruitment Finished |
| 14/NW/0130 | 149373 | A randomised, double blind, multi-center, placebo-controlled study to evaluate the efficacy, safety, and tolerability of NT100 in pregnant women with a history of unexplained recurrent pregnancy loss (RPL) | Number Agreed | 5 | 5 | Not Available / Not Agreed | 17 | 26/10/2015 | Recruitment Finished |
| 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 who w | Number Agreed | 2 | 2 | Not Available / Not Agreed | 3 | 15/09/2015 | Recruitment Finished |
| 14/LO/1192 | 149876 | A Multicenter Phase 3 Randomized Open-Label Study of Bosutinib versus Imatinib in Adult Patients with Newly Diagnosed Chronic Phase Chronic Myelogenous Leukaemia | Number Agreed | 3 | 3 | Not Available / Not Agreed | 3 | 31/08/2015 | Recruitment Finished |
| 15/LO/0034 | 169816 | OPuS-2: A MULTICENTER, RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED, PARALLEL GROUP STUDY TO EVALUATE THE EFFICACY AND SAFETY OF TWO DOSE LEVELS OF BCX4161 FOR 12 WEEKS AS AN ORAL PROPHYLAXIS TREATMENT FOR ATTACKS OF HEREDITARY ANGIOEDEMA | Number Agreed | 3 | 3 | Not Available / Not Agreed | 3 | 15/09/2015 | 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 gabapentin. | 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 |

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| 13/WA/0080 | 120794 | OBSERVATIONAL STUDY PROTOCOL IM101348ST LONGTERM EXPERIENCE WITH ABATACEPT SC IN ROUTINE CLINICAL PRACTICE STUDY | Number Agreed | 6 | 6 | Not Available / Not Agreed | 6 | 20/04/2015 | Recruitment Finished |
| 14/NW/1257 | 161881 | A RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED 2-WAY CROSSOVER STUDY TO EVALUATE THE EFFICACY, SAFETY AND TOLERABILITY OF PF-03715455 ADMINISTERED TWICE DAILY BY INHALATION FOR 4 WEEKS IN SUBJECTS WITH MODERATE TO SEVERE CHRONIC OBSTRUCTIVE PULMONARY DISEASE | Number Agreed | 8 | 8 | Not Available / Not Agreed | 0 | 15/04/2015 | Withdrawn By Sponsor |
| 15/LO/0495 | 173451 | TOLERABILITY OF PF-03715455 ADMINISTERED TWICE DAILY BY INHALATION FOR 4 WEEKS IN SUBJECTS WITH MODERATE TO SEVERE | Number Agreed | 7 | 7 | Not Available / Not Agreed | 0 | 21/07/2015 | Withdrawn By Sponsor |
| 12/SC/0633 | 114037 | CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) | Number Agreed | 3 | 3 | Not Available / Not Agreed | 1 | 20/05/2015 | Recruitment Finished |
| 14/SC/0065 | 145868 | A Randomized, Multicenter, Double-Blind, Parallel, Placebo-Controlled Study of the Effects of Canagliflozin on Renal Endpoints in Adult Subjects With Type 2 Diabetes Mellitus | Number Agreed | 8 | 8 | Not Available / Not Agreed | 5 | 15/03/2015 | Recruitment Finished |
| 14/NW/1053 | 157943 | A Randomized, Double Blind, Placebo-Controlled, Dose Titration, Phase 2 Study to Assess the Safety, Tolerability, Pharmacokinetics, and Pharmacodynamics of ISIS 494372 Administered Subcutaneously to Patients with High Lipoprotein(a) | Number Agreed | 3 | 3 | Not Available / Not Agreed | 3 | 17/06/2015 | Recruitment Finished |
| 15/LO/0032 | 148409 | RANDOMIZED, MULTICENTER, PHASE III, OPEN LABEL STUDY OF ALECTINIB VERSUS CRIZOTINIB IN TREATMENT NA?VE ANAPLASTIC LYMPHOMA KINASE?POSITIVE ADVANCED NON? SMALL CELL LUNG CANCER | Number Agreed | 2 | 2 | Not Available / Not Agreed | 1 | 08/12/2015 | 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, including 28 D | Number Agreed | 3 | 3 | Not Available / Not Agreed | 1 | 06/01/2016 | Recruitment Finished |
| 15/LO/0967 | 178036 | iNNOVATE Study: A randomized, double-blind, placebo-controlled, phase 3 study of ibrutinib or placebo in combination with Rituximab in Subjects with Waldenstroms Macroglobulinemia | Number Agreed | 5 | 5 | Not Available / Not Agreed | 0 | 13/01/2016 | Recruitment Finished |