

**Performance in Initiating and Delivering (PID) Clinical Research**  
**UCLH Performance in Approving and Recruiting to Research**  
**September 2017**

Since July 2012, providers of NHS services have been providing data on performance in initiating and delivering clinical research. Data for studies which have received NHS approval (called Decision to Deliver) in the previous 12 months is submitted quarterly to the National Institute for Health Research (NIHR). Below, a summary of performance submitted by UCLH in Quarter 2 (2017/18) for studies where UCLH was selected as a site over the past year.

Providers are asked to publish details of the length of time taken to approve clinical trials from the point from Date Site Selected to recruitment of the first participant. This period is known as the initiation stage and the NIHR has assigned a benchmark target of 70 days.

Providers also submit information on those Commercially Sponsored Trials, which have closed to recruitment in the year and whether the trial has achieved its anticipated total recruitment target. The period between recruitment of the first participant and reaching the total recruitment target is known as the delivery stage. Each quarter, UCLH will update this information with the most recent studies. The data is correct as of end of September 2017.

For Further information on PID reporting, please visit the NIHR pages at:

<http://www.nihr.ac.uk/policy-and-standards/Performance-in-initiating-and-delivering-research.htm>

Initiating Clinical Research – HRA Approval

| ID | Research Ethics Committee Reference Number | Integrated Research Application System Number | Name of Trial   | Date Site Invited | Date Site Selected | HRA Approval Date | Date Site Confirmed By Sponsor | Date Site Confirmed | Non-Confirmation Status | Date Site Ready To Start | Date of First Patient Recruited | Benchmark Met | Reasons for delay correspond to: |
|----|--|---|---|-------------------|--------------------|-------------------|--------------------------------|---------------------|-------------------------|--------------------------|---------------------------------|---------------|----------------------------------|
| 1  | 16/WM/0437                                 | 206855  | ARIEL4 (Assessment of Rucaparib In Ovarian CancEr Trial): A Phase 3 Multicenter, Randomized Study of Rucaparib versus Chemotherapy in Patients with Relapsed, BRCA-Mutant, High-Grade Epithelial Ovarian, Fallopian Tube, or Primary Peritoneal Cancer  |                   | 05/10/2016         | 15/12/2016        |                                | 26/01/2017          |                         | 08/02/2017               | 28/02/2017                      | No            | Sponsor                          |
| 2  | 16/NE/0342                                 | 213304  | A Phase 1 Dose-escalation Study to Evaluate the Tolerability, Safety, Pharmacokinetics, and Antitumor Activity of ADCT-402 in Patients with Relapsed or Refractory B-cell Lineage Non Hodgkin Lymphoma (B-NHL)  |                   | 11/10/2016         | 24/01/2017        |                                | 02/02/2017          |                         | 15/02/2017               | 05/04/2017                      | No            | Sponsor                          |
| 3  | 16/EE/0243                                 | 207331  | A 6-Month, Multicenter, Phase 3, Open-Label Extension Safety Study Of OTO-104 Given At 3-Month Intervals by Intratympanic Injection in Subjects with Unilateral Meniere's Disease   | 22/09/2016        | 12/10/2016         | 01/08/2016        |                                | 30/11/2016          |                         | 20/12/2016               | 06/02/2017                      | No            | Sponsor                          |
| 4  | 16/SC/0271                                 | 187103  | The UK Plasma based Molecular Profiling of Advanced Breast Cancer to inform Therapeutic Choices (plasmaMATCH) Trial: A multi parallel cohort, open-label, multi-centre phase IIa clinical trial aiming to provide proof of principle efficacy for designated targeted therapies in patients with advanced breast cancer where the targetable mutation is identified through ctDNA screening |                   | 12/10/2016         | 21/09/2016        |                                | 27/01/2017          |                         | 02/02/2017               | 09/03/2017                      | No            | Sponsor                          |

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| 5  | 16/LO/1341 | 191779 | A Phase IIA Prospective, Single-Centre, Open Label Clinical Trial to Evaluate the Safety, Tolerability and Pharmacodynamic Effects of Ambroxol in Patients with Parkinson Disease  | 01/09/2016 | 13/10/2016 | 14/10/2016 |  | 21/11/2016 |  | 23/11/2016 | 11/01/2017 | No | Sponsor |
| 6  | 16/LO/1312 | 188434 | A Phase 2, Randomized, Double-Blind, Placebo-Controlled Trial of IMO-8400-211 in Patients with Dermatomyositis   |            | 13/10/2016 | 02/12/2016 |  | 13/01/2017 |  | 19/01/2017 | 17/05/2017 | No | Sponsor |
| 7  | 16/LO/1612 | 201997 | Prospective Follow-up Study for Patients who Completed Study ALX0681-C301 (HERCULES) to Evaluate Long-term Safety and Efficacy of Caplacizumab (Post-HERCULES)   | 28/09/2016 | 17/10/2016 | 16/11/2016 |  | 14/12/2016 |  | 15/12/2016 | 06/01/2017 | No | Sponsor |
| 8  | 16/LO/1605 | 201385 | HyPer: A Phase 1, Dose Escalation Study of Guadecitabine (SGI-110) a Second Generation Hypo-Methylating Agent in Combination with Pembrolizumab (MK3475) in Patients with Refractory Solid Tumours   |            | 25/10/2016 |            |  | 02/03/2017 |  | 08/03/2017 | 02/05/2017 | No | Sponsor |
| 9  | 16/SC/0304 | 166367 | A Study into the Pharmacodynamic Biomarker Effects of Olaparib (a PARP Inhibitor) ± Degarelix (a GnRH antagonist) given Prior to Radical Prostatectomy   |            | 01/11/2016 | 16/08/2016 |  | 09/02/2017 |  | 10/02/2017 |            | No | Sponsor |
| 10 | 16/LO/1628 | 199315 | The ACL SNNAP Trial: ACL Surgery Necessity in Non Acute Patients. Comparison of the clinical and cost effectiveness of two management strategies for non-acute Anterior Cruciate Ligament (ACL) injury: Rehabilitation versus surgical Reconstruction. |            | 01/11/2016 | 23/08/2016 |  | 21/02/2017 |  | 21/02/2017 |            | No | Sponsor |

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| 11 | 16/LO/1636 | 172808 | A prospective randomized multi-center study comparing endoscopic pneumodilation and per oral endoscopic myotomy (POEM) as treatment of idiopathic achalasia   |            | 03/11/2016 | 03/11/2016 |  | 27/01/2017 |  | 27/01/2017 | 31/03/2017 | No | Sponsor |
| 12 | 16/WM/0406 | 210454 | Red cell transfusion in Acute myeloid Leukaemia   | 01/11/2016 | 10/11/2016 | 08/11/2016 |  | 12/12/2016 |  | 21/12/2016 | 15/06/2017 | No | Neither |
| 13 | 16/EM/0380 | 198726 | A phase II, multicentre, randomised trial comparing combination gemcitabine/carboplatin and hydroxychloroquine versus carboplatin/etoposide therapy alone in small cell lung cancer (SCLC)                              |            | 10/11/2016 | 09/11/2016 |  | 18/01/2017 |  | 20/01/2017 | 26/07/2017 | No | Neither |
| 14 | 16/LO/2008 | 208047 | A Single Arm, Open-Label, Multi-Centre, Phase I/II Study Evaluating the Safety and Clinical Activity of AUTO2, a CAR T Cell Treatment Targeting BCMA and TACI, in Patients with Relapsed or Refractory Multiple Myeloma | 13/06/2016 | 15/11/2016 | 31/03/2017 |  | 03/05/2017 |  | 03/05/2017 | 05/05/2017 | No | Sponsor |
| 15 | 16/LO/1782 | 213379 | Phase Ib/II Trial of Pembrolizumab (MK-3475) Combination Therapies in Metastatic Castration-Resistant Prostate Cancer (mCRPC) (KEYNOTE-365)   |            | 23/11/2016 | 29/11/2016 |  | 08/02/2017 |  | 09/02/2017 | 01/03/2017 | No | Sponsor |
| 16 | 16/LO/1436 | 205613 | Evaluation of Potential Predictors of Disease Progression in Patients with aHUS including Genetics, Biomarkers, and Treatment.  | 14/09/2016 | 29/11/2016 | 29/11/2016 |  | 08/06/2017 |  | 12/06/2017 | 16/06/2017 | No | Sponsor |
| 17 | 16/SC/0418 | 198404 | The clinical effectiveness of 5% Carbamide Plus, 10% Carbamide Plus compared to two commercially available 10% Carbamide peroxide products: a double blind randomised placebo controlled clinical trial                 |            | 07/12/2016 | 07/12/2016 |  | 03/02/2017 |  | 22/02/2017 | 02/05/2017 | No | Sponsor |

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| 18 | 16/LO/2106 | 215913 | A Phase IB Combination Study of Rucaparib (CO-338) and Atezolizumab (MPDL3280A) in Patients with Solid Tumours and advanced gynaecologic cancers, with a focus on Ovarian Cancer                                 | 27/10/2016 | 09/12/2016 | 13/02/2017 |  | 29/03/2017 |  | 03/04/2017 | 07/04/2017 | No | Sponsor |
| 19 | 16/WM/0439 | 211177 | A Multicenter, Randomized, Double-Blind, Sham-Controlled Clinical Investigation of the EndoStim® Lower Esophageal Sphincter (LES) Stimulation System for the Treatment of Gastroesophageal Reflux Disease (GERD) | 17/06/2016 | 12/12/2016 | 12/12/2016 |  | 13/02/2017 |  | 07/04/2017 | 23/05/2017 | No | Neither |
| 20 | 16/SC/0566 | 213579 | A PROSPECTIVE INTERNATIONAL MULTICENTRE RANDOMISED CONTROLLED SINGLE BLIND CLINICAL INVESTIGATION OF MAGNETICALLY ENHANCED DIFFUSION FOR ACUTE ISCHAEMIC STROKE (MEDIS-INTERNATIONAL)                            |            | 15/12/2016 | 10/01/2017 |  | 07/03/2017 |  | 07/03/2017 |            | No | Sponsor |
| 21 | 16/NE/0370 | 214375 | A Phase III Double-blind, Randomized, Placebo-Controlled Study assessing the Efficacy, Safety and Tolerability of Idebenone in Patients with Duchenne Muscular Dystrophy Receiving Glucocorticoid Steroids       |            | 06/01/2017 | 29/12/2016 |  | 14/03/2017 |  | 20/03/2017 | 22/06/2017 | No | Sponsor |
| 22 | 16/LO/1677 | 191232 | A phase II, multi-centre, non-randomised, molecularly stratified trial for NSCLC patients to study tumour heterogeneity using genomic analysis   | 27/10/2016 | 16/01/2017 |            |  | 18/04/2017 |  | 18/04/2017 | 16/05/2017 | No | Sponsor |
| 23 | 16/LO/1502 | 178292 | Utilising Circulating Tumour Cell (CTC) Counts to Optimize Systemic Therapy of Metastatic Prostate Cancer: CTC-STOP Trial  | 16/01/2017 | 16/01/2017 | 07/10/2016 |  | 02/05/2017 |  | 27/04/2017 |            | No | Sponsor |

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| 24 | 16/LO/2186 | 188540 | A randomised phase I/II study of Intensity Modulated Arc Therapy techniques in abdominal neuroblastoma  |            | 17/01/2017 | 17/01/2017 |  | 21/02/2017 |  | 21/02/2017 | 01/03/2017 | Yes |         |
| 25 | 16/LO/1999 | 200065 | A comparison of Cognitive Behaviour Therapy for insomnia (CBTi) and usual audiological rehabilitation in the management of tinnitus related insomnia  |            | 17/01/2017 | 09/02/2017 |  | 24/02/2017 |  | 24/02/2017 | 26/06/2017 | No  | Sponsor |
| 26 | 16/LO/1776 | 191954 | First-in-man feasibility study to assess the safety of TIPS microspheres in perianal fistulas   |            | 18/01/2017 | 17/01/2017 |  | 17/02/2017 |  | 17/02/2017 | 06/06/2017 | No  | Sponsor |
| 27 | 17/SC/0033 | 218114 | A phase 3 randomized, open-label, multicenter study comparing isatuximab (SAR650984) in combination with pomalidomide and low-dose dexamethasone versus pomalidomide and low-dose dexamethasone in patients with refractory or relapsed and refractory multiple myeloma | 11/10/2016 | 19/01/2017 | 29/03/2017 |  | 13/04/2017 |  | 13/04/2017 | 30/05/2017 | No  | Sponsor |
| 28 | 17/LO/0011 | 217376 | VEROnA: A pilot, open label, single-arm, window of opportunity study of vandetanib-eluting radio-opaque embolic beads (BTG 002814) in patients with resectable hepatocellular carcinoma and in patients with resectable liver malignancies                              | 22/11/2016 | 19/01/2017 | 15/02/2017 |  | 08/05/2017 |  | 09/05/2017 | 11/08/2017 | No  | Sponsor |

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| 29 | 16/LO/2148 | 219354 | A Multi-Centre, Open Label, Phase IIb Study, Evaluating the Safety, Tolerability and Efficacy of Targeted Intraprostatic Administration of PRX302 to Treat Men with Histologically Proven, Clinically Significant, Localised, Low- to Intermediate- Risk Prostate Cancer that is Associated with an MRI Lesion | 23/01/2017 | 24/01/2017 |            |  | 28/07/2017 |  | 31/07/2017 | 14/09/2017 | No | Sponsor |
| 30 | 16/LO/2150 | 201093 | Randomised phase II Trial of olaparib, chemotherapy or olaparib and cediranib in patients with BRCA mutated platinum-resistant ovarian cancer  | 25/11/2016 | 27/01/2017 |            |  | 12/05/2017 |  | 12/05/2017 | 25/07/2017 | No | Sponsor |
| 31 | 16/LO/1755 | 209250 | Evaluation of the effect of Duodenal Mucosal Resurfacing (DMR) using the Revita System in the treatment of Type 2 diabetes (T2D)- Revita 2 – C-30000   |            | 31/01/2017 | 24/01/2017 |  | 01/03/2017 |  | 02/03/2017 | 28/04/2017 | No | Neither |
| 32 | 16/NE/0183 | 197040 | Treatment of Poor-grade Subarachnoid Haemorrhage Trial 2   | 08/11/2016 | 31/01/2017 | 13/07/2016 |  | 25/04/2017 |  | 27/04/2017 |            | No | Sponsor |
| 33 | 17/LO/0247 | 213278 | A Phase 1/2, Open-Label, Dose-Escalation, Safety and Tolerability Study of INCAGN01949 in Subjects with Advanced or Metastatic Solid Tumour  | 26/10/2016 | 06/02/2017 | 24/05/2017 |  | 23/08/2017 |  | 24/08/2017 | 18/09/2017 | No | Sponsor |
| 34 | 17/LO/0169 | 214075 | Effect of MD1003 in progressive multiple sclerosis: a randomized double blind placebo controlled study   | 14/11/2016 | 13/02/2017 | 21/04/2017 |  | 09/05/2017 |  | 26/05/2017 | 21/06/2017 | No | Neither |

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| 35 | 17/LO/0284 | 221453 | A study of single doses to evaluate the safety, tolerability, pharmacokinetics and target engagement of nebulised GSK3008348 in idiopathic pulmonary fibrosis patients, using positron emission tomography (PET) imaging  | 21/03/2017 | 15/02/2017 |            |  | 11/05/2017 |  | 12/05/2017 | 13/06/2017 | No | Sponsor |
| 36 | 16/LO/0986 | 202286 | A Randomized, Multicenter, Double Blind, Phase III Study of Nivolumab or Placebo in Subjects with Resected Lower Esophageal, or Gastroesophageal Junction Cancer  | 21/12/2016 | 16/02/2017 | 10/08/2016 |  | 05/04/2017 |  | 12/04/2017 |            | No | Neither |
| 37 | 16/WS/0197 | 186191 | An adaptive multi-arm phase II trial of maintenance targeted therapy after chemotherapy in metastatic urothelial cancer   | 03/02/2017 | 16/02/2017 |            |  | 18/05/2017 |  | 02/06/2017 | 20/09/2017 | No | Sponsor |
| 38 | 16/LO/1004 | 207544 | Efficacy and safety of low-dose IL-2 (ld-IL-2) as a Treg enhancer for anti-inflammatory therapy in newly diagnosed Amyotrophic Lateral Sclerosis (ALS) patients: A randomized, double-blind, placebo-controlled, phase-II Proof of Concept/ Proof of Mechanism Clinical Trial | 06/09/2016 | 20/02/2017 |            |  | 19/06/2017 |  | 22/06/2017 |            | No | Sponsor |
| 39 | 17/LO/0285 | 211245 | Phase 1/2 Safety, Pharmacokinetic, and Antitumor Activity Study of G1T38 in Combination with Fulvestrant in Patients with Hormone Receptor-Positive, HER2 Negative Locally Advanced or Metastatic Breast Cancer after Endocrine Failure                                       | 05/12/2016 | 24/02/2017 | 05/04/2017 |  | 15/05/2017 |  | 25/05/2017 |            | No | Sponsor |



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| 40 | 16/SC/0376 | 202786 | Testing Radical prostatectomy in men with prostate cancer and oligoMetastases to the bone: a randomised controlled feasibility trial  |            | 27/02/2017 | 10/10/2016 |  | 10/03/2017 |  | 13/03/2017 | 19/05/2017 | No | Neither |
| 41 | 17/NE/0058 | 219540 | A Phase 1 Dose Escalation and Cohort Expansion Study of TSR-042, an anti-PD-1 Monoclonal Antibody, in Patients with Advanced Solid Tumors   | 24/02/2017 | 27/02/2017 | 13/04/2017 |  | 14/06/2017 |  | 14/06/2017 | 26/06/2017 | No | Sponsor |
| 42 | 16/SC/0590 | 191279 | DEpletion of Serum amyloid P component In Alzheimer's Disease: DESPIAD. Double-blind placebo controlled randomised phase IIb trial of SAP depletion by CPHPC in mild Alzheimer's disease                                | 12/12/2016 | 02/03/2017 | 16/02/2017 |  | 09/05/2017 |  | 25/05/2017 |            | No | Sponsor |
| 43 | 16/LO/1810 | 209789 | A Phase 3, Randomized Study to Assess the Efficacy and Safety of Ublituximab in Combination with TGR-1202 Compared to Obinutuzumab in Combination with Chlorambucil in Patients with Chronic Lymphocytic Leukemia (CLL) | 10/02/2017 | 02/03/2017 | 03/01/2017 |  | 09/06/2017 |  | 16/06/2017 |            | No | Sponsor |
| 44 | 16/LO/2141 | 200571 | Biological Medicine for Diffuse Intrinsic Pontine Glioma (DIPG) Eradication   | 26/01/2017 | 03/03/2017 | 06/04/2017 |  | 17/05/2017 |  | 29/06/2017 | 07/07/2017 | No | Sponsor |
| 45 | 17/NW/0175 | 222859 | Phase 1b Open-Label, Dose Escalation Study of PRTX-100 in Adult Patients with Persistent/Chronic Immune Thrombocytopenia  | 07/03/2017 | 06/03/2017 |            |  | 27/06/2017 |  | 27/06/2017 | 27/09/2017 | No | Sponsor |

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| 46 | 17/EM/0032 | 209811 | Reclassifying constipation using magnetic resonance imaging combined with high resolution manometry: a validation study and double-blind crossover trial (RECLAIM study)   | 09/12/2016 | 07/03/2017 |            |  | 20/04/2017 |  | 26/04/2017 | No                 | Sponsor |
| 47 | 16/LO/1960 | 199019 | An Open-label, Sequential, Ascending, Repeated Dose-finding Study of Sarilumab, administered with Subcutaneous (SC) Injection, in Children and Adolescents, Aged 1 to 17 years, with Systemic Juvenile Idiopathic Arthritis (sJIA), Followed by an Extension Phase | 07/11/2016 | 07/03/2017 | 28/02/2017 |  | 07/03/2017 |  | 20/04/2017 | No                 | Neither |
| 48 | 17/WM/0105 | 214955 | A Phase 1/2 dose Escalation and Expansion Study to Investigate the safety, pharmacokinetics, pharmacodynamics and clinical activity of GSK525762 in combination with Fulvestrant in subjects with ER+ breast cancer  | 27/10/2016 | 08/03/2017 |            |  | 31/05/2017 |  | 01/06/2017 | No                 | Neither |
| 49 | 17/EE/0205 | 224373 | A Phase I Open-Label Dose Escalation Study to Determine the Efficacy, Safety and Pharmacokinetics of GMI-1271 as Adjunct to Standard of Care Chemotherapy for the Treatment of Multiple Myeloma  | 16/11/2016 | 15/03/2017 |            |  |            |  |            | Site Not Confirmed | Neither |
| 50 | 17/EM/0063 | 213979 | A Phase 3 Multi-Center, 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   | 13/10/2016 | 15/03/2017 | 27/04/2017 |  | 23/05/2017 |  | 02/06/2017 | No                 | Sponsor |

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| 51 | 16/EE/0463 | 214371 | COMPLEMENT-1: An open-label, multicenter, Phase IIIb study to assess the safety and efficacy of ribociclib (LEE011) in combination with letrozole for the treatment of men and postmenopausal women with hormone receptor positive (HR+) HER2-negative (HER2-) advanced breast cancer (aBC) with no prior hormonal therapy for advanced disease |            | 16/03/2017 |            |  | 16/03/2017 |  | 16/03/2017 | 06/04/2017 | Yes |         |
| 52 | 16/SC/0657 | 182046 | A Phase Ib/II combination of acalabrutinib with rituximab, cyclophosphamide, doxorubicin, vincristine and prednisolone (R-CHOP) for patients with Diffuse Large B-cell Lymphoma (DLBCL)   | 20/03/2017 | 22/03/2017 | 28/02/2017 |  | 23/05/2017 |  | 23/05/2017 | 24/07/2017 | No  | Sponsor |
| 53 | 17/YH/0050 | 207335 | An Evaluation of Mindfulness Based Stress Reduction Groups for Adolescents and Young Adults with Crohn's disease and Ulcerative Colitis   | 16/02/2017 | 24/03/2017 | 02/03/2017 |  | 11/04/2017 |  | 12/04/2017 | 25/04/2017 | Yes |         |
| 54 | 16/NE/0279 | 198051 | Risk-stratified sequential Treatment with Ibrutinib and Rituximab (IR) and IR-CHOP for De-novo post-transplant Lymphoproliferative disorder (PTLD)  | 04/01/2017 | 24/03/2017 |            |  | 11/04/2017 |  | 11/04/2017 | 16/05/2017 | Yes |         |
| 55 | 17/WM/0017 | 201600 | A randomised placebo-controlled trial of mifepristone and misoprostol versus misoprostol alone in the medical management of missed miscarriage (MifeMiso trial)   | 16/02/2017 | 11/04/2017 | 06/04/2017 |  | 27/06/2017 |  | 06/07/2017 |            | No  | Sponsor |
| 56 | 17/LO/0475 | 220177 | The Clinical And Biological effects of the use of pRoblOtic VSL#3 in patients with oral lichen planus: a proof-of-concept study (CABRIO)  |            | 12/04/2017 | 12/04/2017 |  | 18/05/2017 |  | 25/05/2017 | 24/08/2017 | No  | Sponsor |

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| 57 | 17/NE/0115 | 218417 | MULTICENTER, INTERNATIONAL, DOUBLE-BLIND, TWO-ARM, RANDOMIZED, PLACEBO-CONTROLLED PHASE II TRIAL OF PIRFENIDONE IN PATIENTS WITH UNCLASSIFIABLE PROGRESSIVE FIBROSING ILD  | 16/01/2017 | 17/04/2017 | 26/05/2017 |  | 23/06/2017 |  | 14/07/2017 |            | No             | Neither |
| 58 | 17/SC/0185 | 214076 | A Phase 3, Randomized, Open-Label, Multicenter Study Comparing the Efficacy and Safety of the Bruton's Tyrosine Kinase (BTK) Inhibitors BGB-3111 and Ibrutinib in Subjects with Waldenström's Macroglobulinemia (WM)                           | 10/11/2016 | 18/04/2017 |            |  | 28/07/2017 |  | 08/08/2017 | 22/08/2017 | No             | Sponsor |
| 59 | 17/LO/0440 | 217506 | Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Safety, Tolerability, Pharmacokinetics and Pharmacodynamics of Multiple Ascending Doses of Intrathecally Administered ISIS 814907 in Patients with Mild Alzheimer's Disease | 26/04/2017 | 26/04/2017 | 20/06/2017 |  | 21/07/2017 |  | 25/07/2017 | 23/08/2017 | No             | Sponsor |
| 60 | 16/WM/0472 | 211270 | CheckPOINT blockade For Inhibition of Relapsed Mesothelioma (CONFIRM): A Phase III Trial to Evaluate the Efficacy of Nivolumab in Relapsed Mesothelioma  | 26/04/2017 | 04/05/2017 | 14/03/2017 |  | 07/06/2017 |  | 01/08/2017 |            | No             | Neither |
| 61 | 16/YH/0157 | 204585 | Personalising Anal cancer radioTherapy dose – Incorporating ACT3, ACT4 and ACT5  | 05/05/2017 | 14/09/2017 |            |  |            |  |            |            | Within 70 days |         |
| 62 | 17/EM/0166 | 222665 | A Phase 2a trial of Avelumab, an anti-PD-L1 antibody, in relapsed and refractory peripheral T-cell lymphoma (PTCL)   | 05/05/2017 | 04/08/2017 |            |  | 07/08/2017 |  | 16/08/2017 |            | Within 70 days |         |

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| 63 | 17/WM/0106 | 201079 | MOTILITY: Small bowel motility quantified by cine MRI as a predictor of long term response in patients with Crohn's disease commencing biological therapy   | 27/01/2017 | 08/05/2017 | 28/04/2017 |  | 02/06/2017 |  | 13/06/2017 | 30/08/2017 | No                 | Sponsor |
| 64 | 17/EM/0155 | 206803 | POINT: Multicenter, randomized, double-blind, parallel-group, add-on, superiority study to compare the efficacy and safety of ponesimod to placebo in subjects with active relapsing multiple sclerosis who are treated with dimethyl fumarate (Tecfidera®)               | 05/05/2017 | 09/05/2017 |            |  |            |  |            |            | Site Not Confirmed | Neither |
| 65 | 17/EM/0183 | 220783 | A randomised Phase III study to assess the clinical efficacy and safety of 100 mg SC Mepolizumab as an add on to maintenance treatment in adults with severe bilateral Nasal Polyposis  | 27/02/2017 | 11/05/2017 |            |  | 16/08/2017 |  | 23/08/2017 |            | No                 | Sponsor |
| 66 | 17/SC/0055 | 219468 | A three-part randomized, double-blind, placebo-controlled study to investigate the efficacy and safety of secukinumab treatment in Juvenile Idiopathic arthritis subtypes of psoriatic and enthesitis-related arthritis   | 09/05/2017 | 16/05/2017 | 27/03/2017 |  | 25/09/2017 |  | 28/09/2017 |            | No                 | Sponsor |
| 67 | 17/LO/0741 | 216587 | Multicenter, randomized, double-blind, placebo-controlled study to assess the efficacy, safety, pharmacokinetics, and pharmacodynamics of GZ/SAR402671 inpatients with early-stage Parkinson's disease carrying a GBA mutation or other prespecified variant (DEME 33640) | 19/12/2016 | 18/05/2017 |            |  |            |  |            |            | Site Not Confirmed | Sponsor |

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|----|------------|--------|--|------------|------------|------------|--|------------|--|------------|------------|--------------------|---------|
| 68 | 16/EE/0546 | 210292 | Barrett's Oesophagus Trial 3 (BEST3): Cluster randomised controlled trial comparing the Cytosponge-TFF3 test with usual care to facilitate the diagnosis of oesophageal pre-cancer in primary care                                 | 16/05/2017 | 19/05/2017 |            |  |            |  |            |            | Site Not Confirmed | Sponsor |
| 69 | 17/NE/0165 | 217768 | AN OPEN-LABEL, SINGLE-ARM STUDY TO EVALUATE THE EFFECTIVENESS AND SAFETY OF OCRELIZUMAB IN PATIENTS WITH EARLY STAGE RELAPSING REMITTING MULTIPLE SCLEROSIS  | 13/10/2016 | 22/05/2017 |            |  |            |  |            |            | Site Not Confirmed | Sponsor |
| 70 | 17/LO/0736 | 225746 | A Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Study to Evaluate the Efficacy and Safety of Intravenously Administered BMS-986168 in Participants with Progressive Supranuclear Palsy with an Open-Label Extension | 27/09/2016 | 22/05/2017 |            |  |            |  |            |            | Site Not Confirmed | Sponsor |
| 71 | 17/LO/0038 | 182633 | A Randomised phase 3 trial of accelerated versus standard BEP chemotherapy for patients with intermediate and poor-risk metastatic germ cell tumours   | 08/03/2017 | 27/06/2017 |            |  |            |  |            |            | Site Not Confirmed | Sponsor |
| 72 | 17/EE/0120 | 220317 | To study the role of nicotinamide riboside in inducing mitochondrial biogenesis  | 08/03/2017 | 23/05/2017 | 23/05/2017 |  |            |  |            |            | Site Not Confirmed | Neither |
| 73 | 17/EM/0152 | 216307 | A Multi-Center, Randomized, 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                 | 17/11/2016 | 23/05/2017 |            |  | 29/09/2017 |  | 29/09/2017 |            | No                 | Sponsor |
| 74 | 17/LO/0769 | 218626 | Face to face structured education group versus audio-visual information in Irritable Bowel Syndrome: a randomised controlled trial   | 12/05/2017 | 25/05/2017 | 12/05/2017 |  | 02/06/2017 |  | 02/06/2017 | 20/06/2017 | Yes                |         |

|    |            |        |  |            |            |  |  |            |  |            |            |                    |         |
|----|------------|--------|--|------------|------------|--|--|------------|--|------------|------------|--------------------|---------|
| 75 | 17/YH/0181 | 227102 | Phase 2 Study of TAK-659 in Patients With Relapsed or Refractory Diffuse Large B-Cell Lymphoma After at Least 2 Prior Lines of Chemotherapy  | 21/04/2017 | 30/05/2017 |  |  | 25/08/2017 |  | 13/09/2017 | 25/09/2017 | No                 | Sponsor |
| 76 | 17/NW/0312 | 211974 | Phase II study of cabozantinib in patients with metastatic gastrointestinal stromal tumor (GIST) who progressed during neoadjuvant, adjuvant or palliative therapy with imatinib and sunitinib   | 10/05/2017 | 01/06/2017 |  |  | 07/08/2017 |  | 09/08/2017 |            | No                 | Sponsor |
| 77 | 17/NW/0317 | 224142 | A feasibility study to evaluate the acceptability of Krio, a food for special medical purposes (FSMP) for use in the ketogenic diet (KD) with regard to product tolerance, compliance and acceptability  | 30/03/2017 | 02/06/2017 |  |  | 14/08/2017 |  | 30/08/2017 |            | No                 | Neither |
| 78 | 17/LO/1043 | 226184 | A Phase 1/2 Study Exploring the Safety, Tolerability, Effect on the Tumor Microenvironment, and Efficacy of Azacitidine in Combination With Pembrolizumab and Epacadostat in Subjects With Advanced Solid Tumors and Previously Treated Stage IIIB or Stage IV Non-Small Cell Lung Cancer and Stage IV Microsatellite-Stable Colorectal Cancer | 14/02/2017 | 06/06/2017 |  |  |            |  |            |            | Site Not Confirmed | Sponsor |
| 79 | 17/ES/0051 | 223060 | A Phase II Open-Label Study of NUC-1031 in Patients with Platinum-Resistant Ovarian Cancer   | 08/06/2017 | 07/08/2017 |  |  | 28/09/2017 |  | 28/09/2017 |            | No                 | Sponsor |

|    |            |        |   |            |            |            |  |            |  |            |            |                    |         |
|----|------------|--------|---|------------|------------|------------|--|------------|--|------------|------------|--------------------|---------|
| 80 | 17/LO/0494 | 224195 | A Phase 3 Single Arm Study Evaluating the Efficacy and Safety of Gene Therapy in Subjects with Transfusion-dependent $\beta$ -Thalassemia, who have a $\beta 0/\beta 0$ Genotype, by Transplantation of Autologous CD34+ Stem Cells Transduced Ex Vivo with a Lentiviral $\beta A$ -T87Q-Globin Vector in Subjects $\geq 50$ Years of Age | 28/03/2017 | 08/06/2017 | 24/05/2017 |  | 28/06/2017 |  | 30/06/2017 | 02/08/2017 | Yes                |         |
| 81 | 17/NW/0330 | 222996 | A Phase 3, Multicenter, Randomized, Double-Blind, Placebo-controlled Study of AG-120 in Previously-treated Subjects with Nonresectable or Metastatic Cholangiocarcinoma with an IDH1 Mutation   | 26/07/2016 | 12/06/2017 |            |  |            |  |            |            | Site Not Confirmed | Sponsor |
| 82 | 16/SC/0584 | 198941 | A randomised phase II double-blinded placebo-controlled trial of intravenous immunoglobulins and rituximab in patients with antibody-associated psychosis (SINAPPS2)  | 12/06/2017 | 12/06/2017 |            |  | 31/08/2017 |  | 25/09/2017 |            | No                 | Neither |
| 83 | 17/EE/0264 | 228153 | A Phase III, Randomized, Multi-Center, Open-Label, Comparative Global Study to Determine the Efficacy of Durvalumab Or Durvalumab and Tremelimumab in Combination With Platinum-Based Chemotherapy for First-Line Treatment in Patients With Metastatic Non Small-Cell Lung Cancer  | 11/05/2017 | 16/06/2017 |            |  | 12/09/2017 |  | 18/09/2017 |            | No                 | Sponsor |
| 84 | 16/LO/1495 | 207629 | Phase III mechanistic, randomised controlled trial of Stopping Perioperative Angiotensin II Converting Enzyme inhibitors and/or receptor blockers in major noncardiac surgery   | 15/03/2017 | 16/06/2017 |            |  | 27/07/2017 |  | 04/08/2017 |            | No                 | Sponsor |



|    |            |        |  |            |            |            |  |            |  |            |            |                    |         |
|----|------------|--------|--|------------|------------|------------|--|------------|--|------------|------------|--------------------|---------|
| 85 | 17/LO/0427 | 199962 | A Phase I/II Trial of Combination Nab-Paclitaxel and Nintedanib or Nab-Paclitaxel and Placebo In Relapsed NSCLC Adenocarcinoma   | 05/04/2017 | 20/06/2017 |            |  |            |  |            |            | Site Not Confirmed | Sponsor |
| 86 | 17/LO/0893 | 211137 | A Phase 1b/2 Trial of Hu5F9-G4 in Combination with Rituximab in Patients with Relapsed/Refractory B-cell Non-Hodgkin's Lymphoma  | 03/02/2017 | 26/06/2017 |            |  |            |  |            |            | Site Not Confirmed | Sponsor |
| 87 | 16/WM/0512 | 218042 | A Phase 1, Open-Label, Randomised, Repeat Dose, Parallel Group Study to Evaluate the Pharmacokinetics, Safety and Tolerability of Ferric Maltol at Three Dosage Levels in Paediatric Subjects aged 10-17 years of age with iron deficiency (with or without anaemia) | 12/01/2017 | 13/01/2017 |            |  | 18/07/2017 |  | 27/07/2017 |            | No                 | Sponsor |
| 88 | 16/NE/0418 | 211187 | A phase III, multicentre randomised controlled trial to determine the efficacy of Robot assisted radical cystectomy (RARC) and intracorporeal urinary diversion by comparison to Open radical cystectomy (ORC) in patients with bladder cancer.                      | 23/09/2016 | 25/01/2017 | 25/01/2017 |  | 17/02/2017 |  | 17/02/2017 | 21/03/2017 | Yes                |         |
| 89 | 17/LO/0048 | 216896 | A Phase I Dose Escalation, Open-Label Clinical Trial Evaluating the Safety and Efficacy of MAGE-A10c796T in Subjects with Stage IIIb or Stage IV Non-Small Cell Lung Cancer (NSCLC)  | 12/01/2017 | 31/03/2017 | 26/04/2017 |  |            |  |            |            | Site Not Confirmed | Sponsor |
| 90 | 17/LO/0001 | 211800 | A Phase 1/2 , Open-Label and Dose-Finding Study of Adeno-Associated Virus (AAV) Serotype 8 (AAV8)-Mediated Gene Transfer of Human Ornithine Transcarbamylase (OTC) in Adults with Late-Onset OTC Deficiency  | 13/09/2016 | 16/02/2017 |            |  | 23/08/2017 |  | 23/08/2017 |            | No                 | Sponsor |

|    |            |        |  |            |            |            |  |            |  |            |            |                    |         |
|----|------------|--------|--|------------|------------|------------|--|------------|--|------------|------------|--------------------|---------|
| 91 | 16/WM/0501 | 185601 | A Phase I trial of WEE1 inhibition with Chemotherapy and Radiotherapy as adjuvant treatment, and a Window of Opportunity trial with Cisplatin in Patients with Head and Neck Cancer  | 28/02/2017 | 01/03/2017 | 28/02/2017 |  | 11/09/2017 |  | 20/09/2017 |            | No                 | Sponsor |
| 92 | 17/LO/0640 | 204303 | A Phase 3, Randomized, Double-Blind, Placebo-Controlled, Crossover Study to Assess the Efficacy and Safety of UX007 in the Treatment of Movement Disorders associated with Glucose Transporter Type 1 Deficiency Syndrome (Glut1 DS) | 20/10/2016 | 04/04/2017 | 04/08/2017 |  | 25/08/2017 |  | 11/09/2017 |            | No                 | Sponsor |
| 93 | 17/EE/0128 | 222308 | A multi-center, open-label trial investigating the efficacy and safety of continued treatment with tisotumab vedotin in patients with solid tumors known to express tissue factor  | 06/02/2017 | 10/04/2017 | 19/05/2017 |  | 07/08/2017 |  | 09/08/2017 | 08/03/2017 | No                 | Sponsor |
| 94 | 17/YH/0076 | 208944 | Controlling and Lowering Blood Pressure with the MobiusHD™ – Defining Efficacy Markers   | 25/08/2016 | 19/06/2017 | 18/05/2017 |  | 22/06/2017 |  | 21/07/2017 |            | No                 | Sponsor |
| 95 | 17/NW/0228 | 222219 | A Randomized, Open-label, Phase 3 Study Comparing Carfilzomib, Dexamethasone, and Daratumumab to Carfilzomib and Dexamethasone for the Treatment of Patients With Relapsed or Refractory Multiple Myeloma                            | 18/04/2017 | 08/06/2017 |            |  |            |  |            |            | Site Not Confirmed | Sponsor |

|     |            |        |   |            |            |            |  |            |  |            |            |                    |              |
|-----|------------|--------|---|------------|------------|------------|--|------------|--|------------|------------|--------------------|--------------|
| 96  | 17/LO/0627 | 218496 | A Three-month, Randomized, Parallel Active Control, Single and Repeat Dose, Dose-escalation Study of the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics and Preliminary Efficacy of VAL-1221 Delivered Intravenously (IV) in Ambulatory and Ventilator-free Patients with Late-Onset GSD-II (Pompe Disease) | 09/01/2017 | 11/05/2017 | 19/06/2017 |  |            |  |            |            | Site Not Confirmed | Sponsor      |
| 97  | 14/LO/2129 | 163516 | Video assisted thoracoscopic lobectomy versus conventional Open Lobectomy for lung cancer, a multi-centre randomised controlled trial with an internal pilot  | 26/04/2017 | 27/06/2017 |            |  | 25/08/2017 |  | 25/08/2017 |            | No                 | Sponsor      |
| 98  | 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 allogeneic stem cell transplantation   | 10/03/2017 | 15/03/2017 |            |  | 17/05/2017 |  | 20/07/2017 | 11/08/2017 | No                 | NHS Provider |
| 99  | 17/LO/0959 | 223644 | An acceptability, feasibility and Pilot trial: Can virtual reality prepare children for ambulatory surgery?   | 30/06/2017 | 01/07/2017 | 30/06/2017 |  | 06/07/2017 |  | 06/07/2017 | 14/07/2017 | Yes                |              |
| 100 | 14/LO/0187 | 143913 | A DOSE-FINDING PHASE 1 STUDY OF TAS-120 IN PATIENTS WITH ADVANCED SOLID TUMORS WITH OR WITHOUT FIBROBLAST GROWTH FACTOR/RECEPTOR (FGF/FGFR)-RELATED ABNORMALITIES FOLLOWED BY A PHASE 2 STUDY IN PATIENTS WITH ADVANCED SOLID TUMORS WITH FGF/FGFR-RELATED ABNORMALITIES  | 10/07/2017 | 04/07/2017 |            |  |            |  |            |            | Site Not Confirmed | Sponsor      |

|     |            |        |  |            |            |            |  |            |            |            |                    |         |
|-----|------------|--------|--|------------|------------|------------|--|------------|------------|------------|--------------------|---------|
| 101 | 17/LO/0812 | 226255 | A Single-arm, Open-label, Multi-centre, Phase I/II Study Evaluating the Safety and Clinical Activity of AUTO3, a CAR T Cell Treatment Targeting CD19 and CD22 Followed by Consolidation with Anti PD1 Antibody in Patients with Relapsed or Refractory Diffuse Large B Cell Lymphoma   | 10/04/2017 | 06/07/2017 |            |  | 25/08/2017 | 05/09/2017 | 07/09/2017 | Yes                |         |
| 102 | 16/LO/2157 | 217001 | EFFICACY, SAFETY, AND TOLERABILITY OF FOSMETPANTOTENATE (RE-024), A PHOSPHOPANTOTHENATE REPLACEMENT THERAPY, IN PATIENTS WITH PANTOTHENATE KINASE-ASSOCIATED NEURODEGENERATION (PKAN)  | 29/09/2016 | 10/07/2017 | 13/07/2017 |  |            |            |            | Site Not Confirmed | Sponsor |
| 103 | 17/NW/0168 | 218752 | Risk-based, response-adapted, Phase II open-label trial of nivolumab + brentuximab vedotin (N + Bv) for children, adolescents, and young adults with relapsed/refractory (R/R) CD30 + classic Hodgkin lymphoma (cHL) after failure of first-line therapy, followed by brentuximab + bendamustine (Bv + B) for participants with a suboptimal response. | 10/02/2017 | 11/07/2017 | 01/06/2017 |  | 09/08/2017 | 11/08/2017 |            | No                 | Neither |
| 104 | 17/LO/0506 | 220370 | A Single Arm, Open-Label, Multi-Centre, Phase I/II Study Evaluating the Safety and Clinical Activity of AUTO3, a CAR T Cell Treatment Targeting CD19 and CD22 in Paediatric and Young Adult Patients with Relapsed Refractory B-cell Acute Lymphoblastic Leukaemia.  | 10/07/2017 | 12/07/2017 | 05/07/2017 |  | 20/09/2017 | 26/09/2017 |            | No                 | Neither |

|     |            |        |  |            |            |            |  |            |  |            |  |                |
|-----|------------|--------|--|------------|------------|------------|--|------------|--|------------|--|----------------|
| 105 | 17/NE/0234 | 228388 | MK3475-629 – a Phase 2 Open-Label, Single arm study to evaluate the safety and efficacy of Pembrolizumab in participants with recurrent or metastatic cutaneous squamous cell carcinoma  | 24/07/2017 | 25/07/2017 |            |  |            |  |            |  | Within 70 days |
| 106 | 17/EE/0291 | 203703 | A Phase III, Randomized, Open-label, Clinical Trial to Compare Pembrolizumab with Brentuximab Vedotin in Subjects with Relapsed or Refractory Classical Hodgkin Lymphoma   | 15/06/2017 | 31/07/2017 |            |  |            |  |            |  | Within 70 days |
| 107 | 17/SC/0276 | 220437 | A prospective randomised control trial of MAKO medial unicndylar knee arthroplasty versus Jig based Oxford unicompartmental knee arthroplasty with navigation control  | 17/07/2017 | 01/08/2017 | 01/08/2017 |  | 04/04/2017 |  | 15/08/2017 |  | Within 70 days |
| 108 | 17/LO/1019 | 223213 | A Phase I open label clinical trial evaluating the safety and anti tumor activity of autologous T cells expressing enhanced TCRs specific for alpha-fetoprotein (AFPc332 T) in HLA-A2 positive subjects with advanced hepatocellular carcinoma (HCC) | 09/02/2017 | 01/08/2017 | 15/09/2017 |  |            |  |            |  | Within 70 days |
| 109 | 17/LO/0797 | 220423 | A multi-center, non-randomized study to evaluate the safety and effectiveness of the Abre venous self-expanding stent system in patients with symptomatic iliofemoral venous outflow obstruction.  | 28/02/2017 | 02/08/2017 | 01/08/2017 |  |            |  |            |  | Within 70 days |
| 110 | 17/NW/0193 | 216411 | IntAct: Intraoperative Fluorescence Angiography to Prevent Anastomotic Leak in Rectal Cancer   | 26/04/2017 | 02/08/2017 |            |  |            |  |            |  | Within 70 days |
| 111 | 16/EE/0357 | 206501 | Opicapone in clinical practice (OPTIPARK)  | 24/10/2016 | 09/08/2017 | 31/10/2016 |  |            |  |            |  | Within 70 days |

|     |            |        |   |            |            |            |  |            |  |            |  |                |
|-----|------------|--------|---|------------|------------|------------|--|------------|--|------------|--|----------------|
| 112 | 17/LO/1055 | 228923 | A phase IV, prospective, randomised single-blind UK multicentre trial of low-dose versus standard dose rituximab for prevention of relapses in acquired TTP   | 06/06/2017 | 10/08/2017 | 01/08/2017 |  | 31/08/2017 |  | 01/09/2017 |  | Within 70 days |
| 113 | 17/LO/1060 | 187783 | RANDOMISED DOUBLE-BLIND EFFICACY AND MECHANISM STUDY OF SUB-SENSORY SACRAL (OPTIMISED) NEUROMODULATION IN ADULTS WITH FAECAL INCONTINENCE   | 10/08/2017 | 10/08/2017 |            |  |            |  |            |  | Within 70 days |
| 114 | 17/ES/0040 | 223878 | A Phase III, Multicentre, Randomised, Placebo-controlled study of Atezolizumab (Anti-PD-L1 Antibody) as Monotherapy and in combination with Platinum-Based chemotherapy in patients with untreated locally advanced or metastatic urothelial carcinoma                    | 07/08/2017 | 11/08/2017 |            |  |            |  |            |  | Within 70 days |
| 115 | 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 allogeneic stem cell transplantation (REACH 3)  | 27/07/2017 | 14/08/2017 |            |  |            |  |            |  | Within 70 days |
| 116 | 17/LO/1273 | 216265 | A Randomized Phase 3 Study of Nivolumab plus Ipilimumab or Nivolumab Combined with Fluorouracil plus Cisplatin versus Fluorouracil plus Cisplatin in Subjects with Unresectable Advanced, Recurrent or Metastatic Previously Untreated Esophageal Squamous Cell Carcinoma |            | 15/08/2017 |            |  |            |  |            |  | Within 70 days |

|     |            |        |  |            |            |  |  |            |  |            |  |                |
|-----|------------|--------|--|------------|------------|--|--|------------|--|------------|--|----------------|
| 117 | 17/LO/1367 | 225049 | A Phase III, open-label, multicentre, randomised study to investigate the efficacy and safety of atezolizumab compared with chemotherapy in patients with treatment-naive advanced or recurrent (Stage IIIB not amenable for multimodality treatment) or metastatic (Stage IV) non-small cell lung cancer who are deemed unsuitable for platinum-containing therapy. | 25/05/2017 | 16/08/2017 |  |  |            |  |            |  | Within 70 days |
| 118 | 17/LO/0117 | 217367 | Immunotherapy for high risk/relapsed CD19+ Acute Lymphoblastic Leukaemia using CAR T cells to target CD19  | 09/08/2017 | 18/08/2017 |  |  |            |  |            |  | Within 70 days |
| 119 | 17/LO/0738 | 213614 | Evaluation of an interdisciplinary performance arts intervention that combines movement and voice to reduce anxiety and depression among those living with the effects of a stroke.  | 31/08/2017 | 01/09/2017 |  |  | 13/09/2017 |  |            |  | Within 70 days |
| 120 | 17/EE/0384 | 218262 | A Prospective, Randomized, Double-Blinded, Placebo-Controlled, Multinational, Multicenter, Parallel-group, Phase III Study to Evaluate the Efficacy and Safety of Apatinib plus Best Supportive Care (BSC) compared to Placebo plus BSC in Patients with Advanced or Metastatic Gastric Cancer (GC)  | 07/02/2017 | 04/09/2017 |  |  |            |  |            |  | Within 70 days |
| 121 | 17/EM/0217 | 200797 | Investigation of SpaCEOAR for Men with High Risk Prostate Cancer treated with HDR Brachytherapy boost and ExterNal Beam Radiotherapy (ICEMAN)  |            | 06/09/2017 |  |  | 12/09/2017 |  | 12/09/2017 |  | Within 70 days |

|     |            |        |   |            |            |  |  |            |  |            |  |                |
|-----|------------|--------|---|------------|------------|--|--|------------|--|------------|--|----------------|
| 122 | 17/LO/1458 | 204582 | An investigation of the parent-observed link between behaviour symptoms and the dietary management of gastrointestinal symptoms in children with autism spectrum disorder.<br>Can gastrointestinal symptoms be managed by dietary intervention and does this bring benefits to children with autism spectrum disorder (ASD) and their families, over and above improvement in gastrointestinal (GI) symptoms? | 22/08/2017 | 06/09/2017 |  |  |            |  |            |  | Within 70 days |
| 123 | 17/LO/0693 | 226490 | An open-label, multi-center long-term safety roll-over study in patients with tuberous sclerosis complex (TSC) and refractory seizures who are judged by the Investigator to benefit from continued treatment with everolimus after completion of Study CRAD001M2304 (EXIST3)   | 27/06/2017 | 19/09/2017 |  |  | 25/09/2017 |  | 27/09/2017 |  | Within 70 days |
| 124 | 16-LO-0994 | 204296 | A phase 1/2 dose-escalation study to evaluate the safety, tolerability, pharmacokinetics and preliminary efficacy of DTX-SPL8783 (a docetaxel (DTX)-dendrimer conjugate) as monotherapy in patients with advanced solid tumours or in combination with nintedanib in patients with non-small cell lung cancer (NSCLC)   |            | 19/09/2017 |  |  |            |  |            |  | #N/A           |
| 125 | 16/YH/0491 | 203984 | Surveillance of invasive Listeria infections in infants under 90 days of age  | 14/09/2017 | 16/09/2017 |  |  |            |  |            |  | Within 70 days |



|     |            |        |   |            |            |            |  |  |  |  |  |                |
|-----|------------|--------|---|------------|------------|------------|--|--|--|--|--|----------------|
| 126 | 17/EE/0016 | 206408 | A Randomised Clinical Trial Of Manual versus trajectory guided Stereoelectroencephalography Electrode Placement   | 16/08/2017 | 25/09/2017 |            |  |  |  |  |  | Within 70 days |
| 127 | 16/EM/0324 | 204032 | A Phase III Randomized, Placebo-Controlled Clinical Trial to Study the Safety and Efficacy of Suvorexant (MK-4305) for the Treatment of Insomnia in Subjects with Alzheimer's Disease | 20/07/2017 | 25/09/2017 |            |  |  |  |  |  | Within 70 days |
| 128 | 17/ES/0107 | 231506 | A Phase 2 single arm study of Safety and Efficacy of Coversin in adult aHUS subjects  | 20/07/2017 | 26/09/2017 | 14/09/2017 |  |  |  |  |  | Within 70 days |

## Delivering Clinical Research

| ID | 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 | Total Number Of Study Participants Recruited | Reason For Closure Of Trial |
|----|--|---|---|-----------------------------------|---|---|---|--|--|---|--|-----------------------------|
| 1  | 15/EE/0460                                 | 193953  | Evaluation of a Novel Electronic Transanal Irrigation System - Navina Smart | Number Agreed                     | 16  | 16  | Not Available / Not Agreed              |  |  | 01/10/2016                                | 16   | Recruitment Finished        |

|   |            |        |   |               |    |    |                            |            |   |            |    |                      |
|---|------------|--------|---|---------------|----|----|----------------------------|------------|---|------------|----|----------------------|
| 2 | 14/SC/1167 | 144247 | Phase 3, Randomised, PlaceboControlled, Doubleblind, MultiCentre, TwoPart Study of Patritumab (U31287) in Combination with Erlotinib in EGFR Wildtype Subjects with Locally Advanced or Metastatic Non-Small Cell Lung Cancer (NSCLC) Who Have Progressed on at | Number Agreed | 5  | 5  | Date Agreed                | 31/10/2014 | 2 | 06/11/2016 | 2  | Withdrawn By Sponsor |
| 3 | 14/NE/1175 | 164135 | Open Label, Adaptive Design, Ascending, Multiple-Dose Study to Evaluate Safety and Efficacy of BMS-986004 (Anti-CD40L dAb) in Adult Subjects with Primary Immune Thrombocytopenia (ITP)   | Number Agreed | 2  | 2  | Date Agreed                | 01/06/2016 | 0 | 08/11/2016 | 0  | Withdrawn By Sponsor |
| 4 | 13/WM/0373 | 135259 | A Phase 3 Switchover Study of the Efficacy and Safety of BMN 701 (GILT-tagged Recombinant Human GAA) and Long-Term Study for Extended Treatment in rhGAA Exposed Subjects with Late-onset Pompe Disease   | Number Agreed | 3  | 3  | Date Agreed                | 01/06/2015 | 1 | 24/11/2016 | 1  | Withdrawn By Sponsor |
| 5 | 16/LO/0951 | 203461 | A PHASE II PILOT STUDY TO EVALUATE THE SAFETY, TOLERABILITY, EFFICACY, PHARMACODYNAMICS AND PHARMACOKINETICS OF IDES IN ASYMPTOMATIC ANTIBODY-MEDIATED THROMBOTIC THROMBOCYTOPENIC PURPURA (TTP) PATIENTS WITH LOW ADAMTS13 ACTIVITY                            | Number Agreed | 2  | 2  | Not Available / Not Agreed |            |   | 27/12/2016 | 2  | Withdrawn By Sponsor |
| 6 | 15/LO/0681 | 174407 | A Randomized, Double-blind, Placebo-controlled Study to Evaluate the Safety, Tolerability and Pharmacokinetics of Multiple Ascending Doses of Intrathecally Administered ISIS 443139 in Patients with Early Manifest Huntington's Disease                       | Number Agreed | 6  | 6  | Date Agreed                | 01/11/2015 | 6 | 22/02/2017 | 6  | Recruitment Finished |
| 7 | 15/LO/0009 | 146896 | A Prospective Development Study evaluating Focal Therapy using Encage? Coiled Bipolar Radiofrequency Ablation in Men with Localised Prostate Cancer   | Number Agreed | 20 | 20 | Not Available / Not Agreed |            |   | 28/02/2017 | 23 | Recruitment Finished |
| 8 | 15/LO/1500 | 187068 | Investigation of drug-drug interaction between nintedanib and pirfenidone in patients with IPF (an open label, multiple-dose, two group study)  | Number Agreed | 2  | 2  | Date Agreed                | 01/07/2016 | 3 | 14/03/2017 | 3  | Recruitment Finished |

|    |            |        |  |               |    |    |                            |            |    |            |    |                      |
|----|------------|--------|--|---------------|----|----|----------------------------|------------|----|------------|----|----------------------|
| 9  | 14/SC/1340 | 17797  | A Multicenter, Multinational, Randomized, Double Blind, Placebo Controlled, Parallel Group Study to Evaluate the Efficacy and Safety of Laquinimod (0.5, 1.0 and 1.5 mg/day) as Treatment in Patients with Huntington's Disease      | Number Agreed | 10 | 10 | Date Agreed                | 27/11/2015 | 10 | 05/04/2017 | 10 | Recruitment Finished |
| 10 | 16/EM/0408 | 211983 | A Phase 2, Open-Label, Randomized Study to Evaluate the Efficacy and Safety of GS-5745 Combined with Nivolumab versus Nivolumab Alone in Subjects with Unresectable or Recurrent Gastric or Gastroesophageal Junction Adenocarcinoma | Number Agreed | 3  | 3  | Not Available / Not Agreed |            |    | 10/04/2017 | 8  | Recruitment Finished |
| 11 | 14/SC/1340 | 151325 | A Multicenter, Multinational, Randomized, Double Blind, Placebo Controlled, Parallel Group Study to Evaluate the Efficacy and Safety of Laquinimod (0.5, 1.0 and 1.5 mg/day) as Treatment in Patients with Huntington's Disease      | Number Agreed | 7  | 7  | Date Agreed                | 27/11/2015 | 10 | 21/04/2017 | 10 | Recruitment Finished |
| 12 | 13/EM/0373 | 15394  | An Open-label, Multicenter, Phase 2 Study of Oral MLN9708 in Adult Patients With Relapsed and/or Refractory Follicular Lymphoma  | Number Agreed | 3  | 3  | Date Agreed                | 01/01/2015 | 1  | 25/04/2017 | 1  | Recruitment Finished |
| 13 | 15/LO/1441 | 186018 | A Phase III Double-Blind, Randomized, Parallel Group, Multicenter Placebo-Controlled Trial to Study the Efficacy and Safety of Caplacizumab in Patients with Acquired Thrombotic Thrombocytopenic Purpura                            | Number Agreed | 18 | 18 | Date Agreed                | 01/11/2017 | 18 | 29/04/2017 | 18 | Recruitment Finished |
| 14 | 15/LO/1441 | 186018 | A Phase III Double-Blind, Randomized, Parallel Group, Multicenter Placebo-Controlled Trial to Study the Efficacy and Safety of Caplacizumab in Patients with Acquired Thrombotic Thrombocytopenic Purpura                            | Number Agreed | 3  | 3  | Not Available / Not Agreed |            |    | 29/04/2017 | 18 | Recruitment Finished |
| 15 | 15/EM/0454 | 184386 | An International, Multicenter, Prospective, Post Market Registry Using a New Device for Endoscopic Resection of Early Neoplasia in Barrett's Esophagus   | Number Agreed | 24 | 24 | Date Agreed                | 01/06/2016 | 24 | 04/05/2017 | 24 | Recruitment Finished |

|    |            |        |  |               |   |   |                            |            |    |            |    |                      |
|----|------------|--------|--|---------------|---|---|----------------------------|------------|----|------------|----|----------------------|
| 16 | 16/LO/0334 | 193988 | A Phase 3 Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Efficacy and Safety of GS-5745 Combined with mFOLFOX6 as First Line Treatment in Patients with Advanced Gastric or Gastroesophageal Junction Adenocarcinoma                         | Number Agreed | 2 | 2 | Date Agreed                | 01/05/2017 | 2  | 05/05/2017 | 2  | Recruitment Finished |
| 17 | 16/NE/0027 | 187317 | A phase 3, multicenter, randomized, open-label study of avelumab (MSB0010718C) alone or in combination with Pegylated Liposomal Doxorubicin versus Pegylated Liposomal Doxorubicin alone in patients with platinum resistant/refractory ovarian cancer           | Number Agreed | 3 | 3 | Date Agreed                | 01/11/2016 | 23 | 15/05/2017 | 23 | Recruitment Finished |
| 18 | 15/LO/0443 | 172123 | A PHASE 3, MULTICENTER,RANDOMIZED, OPENLABEL STUDY TO COMPARE THE EFFICACY AND SAFETY OF POMALIDOMIDE, BORTEZOMIB AND LOW-DOSE DEXAMETHASONE VERSUS BORTEZOMIB AND LOW-DOSE DEXAMETHASONE IN SUBJECTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA               | Number Agreed | 6 | 6 | Not Available / Not Agreed |            |    | 15/05/2017 | 5  | Recruitment Finished |
| 19 | 14/LO/1597 | 158625 | A Randomized, Double-Blind, Placebo-Controlled, Four-Arm,Parallel-Group, Proof of Concept, and Dose-Finding Adaptive Phase 2a/2b Study to Investigate the Safety, Tolerability and Efficacy and Effect on Quality of Life of Human Recombinant Alkaline Phospha  | Number Agreed | 5 | 5 | Date Agreed                | 01/01/2016 | 12 | 18/05/2017 | 12 | Recruitment Finished |
| 20 | 15/WM/0457 | 195511 | An Open-Label Treatment Use Protocol for Daratumumab in Subjects with Multiple Myeloma Who Have Received at Least 3 Prior Lines of Therapy (Including a Proteasome Inhibitor and an Immunomodulatory Agent) or are Double Refractory to a Proteasome Inhibitor a | Number Agreed | 5 | 5 | Date Agreed                | 01/09/2016 | 7  | 18/05/2017 | 7  | Recruitment Finished |
| 21 | 16/LO/1450 | 202124 | A Phase 3, Multicenter, Randomized, Double Blind Study of Bortezomib and Dexamethasone in Combination with Either Venetoclax or Placebo in Subjects with Relapsed or Refractory Multiple Myeloma Who are Sensitive or Na?ve to Proteasome Inhibitors             | Number Agreed | 4 | 4 | Date Agreed                | 30/06/2018 | 5  | 19/05/2017 | 5  | Recruitment Finished |

|    |            |        |   |               |    |    |                            |            |   |            |    |                      |
|----|------------|--------|---|---------------|----|----|----------------------------|------------|---|------------|----|----------------------|
| 22 | 15/SW/0220 | 182750 | A Phase 3, Multicentre, Randomised, Double-blind, Placebo-controlled Study Evaluating the Efficacy and Safety of Ustekinumab in the Treatment of Subjects With Active Axial Nonradiographic Spondyloarthritis   | Number Agreed | 3  | 3  | Date Agreed                | 17/07/2017 | 1 | 02/07/2017 | 1  | Withdrawn By Sponsor |
| 23 | 16/SW/0090 | 199970 | Ustekinumab in Subjects with Radiographic Axial Spondyloarthritis   | Number Agreed | 3  | 3  | Date Agreed                | 31/07/2017 | 1 | 30/05/2017 | 1  | Withdrawn By Sponsor |
| 24 | 16/SC/0341 | 204761 | A Phase 2, Multicenter Study of the EZH2 Inhibitor Tazemetostat in Adult Subjects with Relapsed or Refractory Malignant Mesothelioma with BAP1 Loss of Function   | Number Agreed | 3  | 3  | Date Agreed                | 01/10/2017 | 2 | 02/06/2017 | 2  | Recruitment Finished |
| 25 | 16/LO/1612 | 201997 | Prospective Follow-up Study for Patients who Completed Study ALX0681-C301 (HERCULES) to Evaluate Long-term Safety and Efficacy of Caplacizumab (Post-HERCULES)  | Number Agreed | 20 | 20 | Not Available / Not Agreed |            |   | 09/06/2017 | 17 | Recruitment Finished |
| 26 | 16/LO/0083 | 193141 | A Phase 3, double-blind, randomized, placebo-controlled, multicenter study to determine the efficacy and safety of luspatercept (ACE-536) plus best supportive care versus best supportive care in adults who require regular red-blood cell transfusions due t | Number Agreed | 3  | 3  | Date Agreed                | 01/02/2018 | 5 | 10/06/2017 | 5  | Recruitment Finished |
| 27 | 15/LO/1670 | 188788 | Randomized, Placebo-Controlled, Double-Blind Phase 2 Study of Patritumab (U3-1287) in Combination with Cetuximab plus Platinum-Based Therapy in First Line Setting in Subjects with Recurrent or Metastatic Squamous Cell Carcinoma of the Head and Neck        | Number Agreed | 12 | 12 | Date Agreed                | 01/04/2017 | 2 | 21/06/2017 | 4  | Recruitment Finished |
| 28 | 16/NW/0379 | 200426 | LY2801653 and LY3009806 - Advanced or Metastatic Biliary Tract Cancer   | Number Agreed | 8  | 8  | Date Agreed                | 01/05/2017 | 4 | 03/07/2017 | 4  | Recruitment Finished |
| 29 | 16/EE/0011 | 194393 | A PROSPECTIVE, RANDOMIZED, DOUBLE BLIND, PLACEBO-CONTROLLED, MULTICENTER, PHASE 3 EFFICACY AND SAFETY STUDY OF OTO-104 GIVEN AS A SINGLE INTRATYMPANIC INJECTION IN SUBJECTS WITH UNILATERAL MENIERE'S DISEASE  | Number Agreed | 3  | 3  | Not Available / Not Agreed |            |   | 24/07/2017 | 1  | Withdrawn By Sponsor |

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|----|------------|--------|---|---------------|----|----|----------------------------|------------|---|------------|----|----------------------|
| 30 | 16/EM/0165 | 203281 | A Phase 3 Multicenter Open-label Study of Brigatinib (AP26113) versus Crizotinib in Patients with ALK-positive Advanced Lung Cancer (CANC 5545)   | Number Agreed | 2  | 2  | Date Agreed                | 22/03/2018 | 2 | 28/07/2017 | 2  | Recruitment Finished |
| 31 | 14/LO/2156 | 166004 | A Phase 2 Proof of Concept (PoC), DoubleBlind, Randomised, Placebocontrolled Study to Evaluate the Efficacy, Safety, Pharmacokinetics and Pharmacodynamics of VSN16R for the treatment of spasticity in patients with Multiple Sclerosis. | Number Agreed | 75 | 75 | Not Available / Not Agreed |            |   | 31/07/2017 | 81 | Recruitment Finished |
| 32 | 16/EE/0243 | 207331 | A 6-Month, Multicenter, Phase 3, Open-Label Extension Safety Study Of OTO-104 Given At 3-Month Intervals by Intratympanic Injection in Subjects with Unilateral Meniere's Disease   | Number Agreed | 5  | 5  | Date Agreed                | 31/03/2017 | 1 | 05/09/2017 | 1  | Withdrawn By Sponsor |
| 33 | 16/SC/0076 | 199027 | A Pilot Double-blind, Placebo-controlled Crossover Study to Explore the Possible Benefit of AUT00063, an Oral Modulator of Voltage-gated Potassium Channels, in Adult Post-lingual Unilateral Cochlear Implant                            | Number Agreed | 8  | 8  | Not Available / Not Agreed |            |   | 27/03/2017 | 2  | Recruitment Finished |