PERFORMANCE IN INITIATING CLINICAL RESEARCH (Q1, 2013/14)

Research Ethics Committee Reference Number	Name of Trial	Date of Receipt of Valid Research Application	Date of First Patient Recruited	Benchmark Met?	Reason for not meeting benchmark
12/SC/0435	Efficacy and safety of 3 doses of S38093 (2, 5 and 20 mg/day) versus placebo, in co-administration with donepezil (10mg/day) in patients with moderate Alzheimer's Disease	19/10/2012	20/05/2013	No	No patients were consented during the 70-day period
GTAC192	Stem Cells in Rapidly Evolving Active Multiple Sclerosis (STREAMS)	19/10/2012	23/03/2013	No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process
11/LO/1465	MAPPING: Diagnostic accuracy of MRI, diffusion-weighted MRI, FDG-PET/CT and FEC-PET/CT in the detection of lymph node metastases in surgically staged endometrial and cervical carcinoma	27/09/2011	19/03/2013	No	Delays during NHS permissions and approvals process
09/S0703/98	TOUCAN (Bladder cancer) Vandetanib in non-cisplatin fit patients with urothelial cancers - A randomised phase II Trial of carboplatin and gemcitabine +/- vandetanib in first line treatment Of advanced Urothelial cell Cancer in patients who are not?	06/01/2011		No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process
12/EE/0165	A Phase 3, Multicenter, Randomized, Double-blind, Active-controlled, Parallel-group Trial with an Open-label Extension Phase to Evaluate the Efficacy and Safety of Oral E5501 versus Eltrombopag, in Adults with Chronic Immune Thrombocytopenia (ITP)	27/04/2012		No	Delays during NHS permissions and approvals process
11/SC/0408	Surgical and large bore pleural procedures in malignant pleural Mesothelioma And Radiotherapy Trial (SMART trial): a randomised controlled trial evaluating whether prophylactic radiotherapy reduces the incidence of procedure tract metastases	01/06/2012		No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process
12/LO/0355	192024-041D - A Multicenter, Paired-eye Comparison, Dose-escalation, Single Dose, 24-Month Study of Safety and Efficacy of Bimatoprost Preservative Free Intracameral Drug Delivery System (Bimatoprost PF IC DDS) in Patients With Open-Angle Glaucoma	18/06/2012		No	Delays during NHS permissions and approvals process
11/SC/0454	A phase IIIb parallel group, open label study of pegylated interferon alfa2a monotherapy (PEGIFN, Ro 258310) compared to untreated control in children with HBeAg positive chronic hepatitis B	08/10/2012	04/04/2013	No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process (sponsor amendments)
11/NW/0548	A Randomised Investigation of Alternative Ofatumumab-containing regimens in less fit patients with CLL (RIAltO version 1)	24/06/2012		No	No eligible patients were identified in clinics during the 70-day period
12/NE/0266	A phase 3 multicenter, randomised, double-blind, placebo-controlled study to investigate the efficacy and safety of GS-7977 + Ribavirin for 12 weeks in treatment naive and treatment experienced subjects with chronic genotype 2 or 3 HCV infection	09/10/2012	29/10/2012	Yes	First patient recruited within 70 day benchmark
11/LO/0958	FASTForward: a randomised clinical trial testing a 1-week course of curative whole breast radiotherapy against a standard 3-week schedule in terms of local cancer control and late adverse effects in women with early breast cancer	28/08/2012	19/09/2012	Yes	First patient recruited within 70 day benchmark
10/MRE00/12	Efficacy of Metformin in Pregnant Obese Women, a Randomised Controlled Trial (EMPOWaR)	24/06/2012		No	Delays during NHS permissions and approvals process

11/LO/1385	A phase 1 study to assess the safety, tolerability and pharmacokinetic profile of boceprevir and sildenafil when dosed separately and together in healthy male volunteers (the boceprevir and sildenafil PK study)	15/11/2012	15/12/2012	Yes	First patient recruited within 70 day benchmark
10/H0402/77	Dexamethasone Reduces Emesis After Major gastrointestinal Surgery (DREAMS trial) - a prospective, doubleblind, multicentre, randomised control trial	27/08/2012	11/10/2012	Yes	First patient recruited within 70 day benchmark
12/LO/0750	Imaging baseline mu opioid receptor availability and stimulated release of endogenous opioids in pathological gamblers using [11C] carfentanil PET and amphetamine challenge	14/08/2012	28/09/2012	Yes	First patient recruited within 70 day benchmark
11/NE/0228	ION Is ablative radiOiodine Necessary for low risk differentiated thyroid cancer patients	09/08/2012	18/09/2012	Yes	First patient recruited within 70 day benchmark
12/YH/0159	HERO: Hydroxychloroquine Effectiveness in Reducing Symptoms of hand OA, a randomised, doubleblind, placebo-controlled trial	11/10/2012	27/11/2012	Yes	First patient recruited within 70 day benchmark
12/LO/1427	Is digital breast tomosynthesis effective in reducing the recall rate in the prevalent screen of women invited to the breast screening programme?	22/11/2012	21/01/2013	Yes	First patient recruited within 70 day benchmark
11/YH/0442	Enhanced Control of Hypertension and Thrombolysis Stroke Trial (ENCHANTED)	05/09/2012	08/11/2012	Yes	First patient recruited within 70 day benchmark
12/LO/1173	An Open-Label Phase I/II Study of GSK2110183 in Combination with Carboplatin and Paclitaxel in Subjects with Platinum-Resistant Ovarian Cancer	20/07/2012	04/02/2013	No	No eligible patients were identified in clinics during the 70-day period
11/LO/1909	Impact of renal denervation on symptomatology, chemoreflex, baroreflux, cardiopulmonary exercise physiology and cardiac performance in patients with chronic heart failure compared with sham procedure	07/08/2012	06/09/2012	Yes	First patient recruited within 70 day benchmark
11/WS/0012	Reducing with Metformin vascular lesions in T1DM (The REMOVAL Study)	10/10/2012	01/12/2012	Yes	First patient recruited within 70 day benchmark
12/LO/0597	Yoga and Cardiovascular Health Trial (YACHT)	26/07/2012	03/10/2012	Yes	First patient recruited within 70 day benchmark
12/LO/1598	A Study to Evaluate the Safety and Effect of ABT-450, Ritonavir and ABT-267 (ABT-450/r/ABT-267) and ABT-333 Coadministered With Ribavirin (RBV) in Hepatitis C Virus (HCV) Genotype 1-infected Adults With Compensated Cirrhosis (TURQUOISE-II)	12/11/2012	28/01/2013	No	Delays during NHS permissions and approvals process
11/SC/0372	ChemoCentryx CL005_140 Study	18/07/2012	03/10/2012	No	No eligible patients were identified in clinics during the 70-day period
12/EE/0176	Randomised, phase IV, placebo-controlled, comparative study to evaluate the efficacy and safety of tapering methotrexate (MTX) dosage versus maintaining the dosage in patients with severe active rheumatoid arthritis (RA)	12/12/2012	28/02/2013	No	No eligible patients were identified in clinics during the 70-day period
12/LO/1597	A Randomized, Double-blind, Placebo-controlled Study to Evaluate the Efficacy and Safety of ABT-450/Ritonavir/ABT-267 (ABT-450/r/ABT-267) and ABT-333 Co-administered with Ribavirin (RBV) in Treatment-Experienced Adults with Genotype 1?(SAPPHIRE-II)	13/11/2012	05/02/2013	No	Delays during NHS permissions and approvals process
12/YH/0318	A Phase 3, Randomized, Double-Blind, Placebo-Controlled Study Evaluating the Efficacy and Safety of GS-1101 (CAL-101) in Combination with Bendamustine and Rituximab for Previously Treated Chronic Lymphocytic Leukemia	01/11/2012	11/01/2013	No	No eligible patients were identified in clinics during the 70-day period

11/LO/0097	VIP: A prospective, phase II, double blinded, multicentre, randomised clinical trial comparing combination gemcitabine and vandetanib therapy with gemcitabine therapy alone in locally advanced or metastatic pancreatic carcinoma	26/06/2012	29/10/2012	No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process
12/YH/0178	A Phase 3, Randomized, Double-Blind, Placebo-Controlled Study Evaluating the Efficacy and Safety of GS-1101 (CAL-101) in Combination with Rituximab for Previously Treated Chronic Lymphocytic Leukemia	13/10/2012	06/02/2013	No	No eligible patients were identified in clinics during the 70-day period
12/NW/0431	A Multi-Centre, Open-Label, Adaptive, Randomised Study of Palifosfamide-tris, a Novel DNA Crosslinker, in Combination with Carboplatin and Etoposide (PaCE) Chemotherapy versus Carboplating and Etoposide (CE) Alone in Chemotherapy Naive Patients?	10/10/2012	26/02/2013	No	Delays during NHS permissions and approvals process
12/NW/0251	A multi-centre, stratified, open, randomized, comparator-controlled, parallel-group phase III study comparing treatment with 177Lu-DOTA0-Tyr3-Octreotate to Octreotide LAR in patients with inoperable, progressive, somatostatin receptor positive?	29/09/2012	08/02/2013	No	No eligible patients were identified in clinics during the 70-day period
12/NW/0045	A RandoMised study of best Available therapy versus JAK Inhibition in patients with high risk Polycythaemia Vera or Essential Thrombocythaemia who are resistant or intolerant to HydroxyCarbamide (MAJIC)	22/06/2012	01/11/2012	No	No eligible patients were identified in clinics during the 70-day period
12/SS/0045	A phase 2, multicentre, randomised, double-blind, placebo-controlled trial of AMG 479 or placebo in combination with gemcitabine as first-line therapy for locally advanced unresectable adenocarcinoma of the pancreas	09/07/2012		No	Sponsor withdrew study before participants could be recruited
12/WA/0051	A stratified phase II study of neoadjuvant chemotherapy given before SCPRT as treatment for patients with MRI-staged operable rectal cancer at high risk of metastatic relapse	24/07/2012	15/01/2013	No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process
12/LO/0570	Randomised clinical trial comparing VNUSClosureFAST with Clarivein for varicose veins	23/06/2012	22/01/2013	No	No eligible patients were identified in clinics during the 70-day period
12/LO/0477	A Phase II study to assess the safety and efficacy of the steroid sulfatase inhibitor Irosustat when added to an aromatase inhibitor in ER positive locally advanced or metastatic breast cancer patients (IRIS)	06/06/2012	06/02/2013	No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process
12/EE/0005	A multicentre, double blind, randomised controlled Clinical Investigation to validate the EPS1 device as a treatment for stroke induced dysphagia - A Study of Swallowing Treatment using Electrical Pharyngeal Stimulation (STEPS Study)	21/06/2012	28/02/2013	No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process
10/MRE00/55	BIG 307 - DCIS (Ductal Carcinoma in situ) Boost Trial	21/01/2011	20/11/2012	No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process
11/NW/0690	Bardoxolone Methyl Evaluation in Patients with Chronic Kidney Disease and Type 2 Diabetes: the Occurrence of Renal Events (BEACON)	01/08/2012		First patient recruited - awaiting confirmation of date	
12/LO/0098	A randomized, double-blind, placebo-controlled, multiple dose study to evaluate the safety, tolerability and efficacy of intravenous administration of secukinumab (AIN457) in patients with asthma not adequately controlled with inhaled?	30/08/2012	13/12/2012	No	No eligible patients were identified in clinics during the 70-day period
12/EE/0055	Research Project in Association with Protocol WN25 203 "CSF sample retention for biomarker assay development"	08/08/2012		No	Delays during NHS permissions and approvals process

10/H1306/5	Phase III Open, Multicentre Study to Investigate the Safety and Efficacy of BPLs High Purity FACTOR X in the Treatment of Factor X Deficient Subjects Undergoing Surgery	02/09/2012		First patient recruited - awaiting confirmation of date	
10/H1005/6	Does co-careldopa treatment in combination with routine NHS occupational and physical therapy, delivered early after stroke within a stroke rehabilitation service, improve functional recovery including walking ability and arm function?	04/09/2012		No	No eligible patients were identified in clinics during the 70-day period
11/LO/1966	A Study to Measure In vivo Changes in Oestrogen Receptor DNA Binding Events in Breast Cancer Treated with Endocrine Therapy for Primary or Recurrent Disease	21/09/2012		First patient recruited - awaiting confirmation of date	
12/WS/0180	A multi-center, two stage, phase II study, evaluating the efficacy of oral BEZ235 plus best supportive care (BSC) versus placebo plus BSC in the treatment of patients with advanced pancreatic neuroendocrine tumors (pNET) after failure of mTOR?	05/10/2012		No	Delays during NHS permissions and approvals process
09/H0406/106	Ofatumumab versus Rituximab Salvage Chemoimmunotherapy followed by ASCT in Relapsed or Refractory DLBCL	10/10/2012		No	Delays during NHS permissions and approvals process
12/LO/1211	A phase III, open-label, multi centre pilot study to assess the feasibility of switching, individuals receiving Atripla or Kivexa plus Efavarinz with continuing Central Nervous System (CNS) toxicity, to a fixed dose of tenofovir/emtricitabine	10/10/2012		First patient recruited - awaiting confirmation of date	
11/NW/0075	Phase III study comparing post-operative conformal radiotherapy to no post-operative radiotherapy in patients with completely resected non-small cell lung cancer and mediastinal nodal N2 involvement	10/10/2012		No	Delays during NHS permissions and approvals process
12/LO/0517	Cognitive pattern and progression in cognitive variant of multiple sclerosis: a follow-up study	10/10/2012		No	No eligible patients were identified in clinics during the 70-day period
12/YH/0179	A Companion Trial to Study GS-US-312-0116: A Phase 3, Randomized, Double-Blind, Placebo-Controlled Study Evaluating the Efficacy and Safety of GS-1101 (CAL-101) in Combination with Rituximab as Therapy for Patients with Previously Treated Chronic?	13/10/2012		No	No eligible patients were identified in clinics during the 70-day period
11/WM/0287	A randomized multi-center study to compare the efficacy of additional tumor debulking surgery versus chemotherapy alone for recurrent platinum sensitive ovarian cancer AGO-OVAR-ID: AGO-OVAR OP.4 (DESKTOP III)	10/10/2012	25/05/2013	No	No patients were consented during the 70-day period / delays during NHS permissions and approvals process
12/LO/0066	A phase II double blind, randomised controlled trial of vegf inhibitor axitinib monotherapy with early dynamic conrast enhanced ultrasound monitoring in chemorefractory third line metastatic colorectal cancer (AXMUS-C)	05/09/2012	17/09/2012	Yes	First patient recruited within 70 day benchmark
12/LO/0926	Ward Randomised Case-crossever trial of a new Personal Hand Hygiene Device by ORBEL	19/10/2012	20/02/2013	No	No eligible patients were identified in clinics during the 70-day period
12/LO/1100	A phase 1 open label, dose-escalation study to asess the safety, pharmacokinetics and activity of NUC-1031, an RRVeoside analogue, in patients with solid advanced tumours	15/08/2012	01/10/2012	Yes	First patient recruited within 70 day benchmark
11/SW/0307	The effect of exercise on wound healing in venous leg ulcer	22/10/2012		No	No eligible patients were identified in clinics during the 70-day period

12/LO/1857	Randomised Phase II study of cetuximab alone or in combination with irinotecan in patients with metastatic CRC with either KRAS WT or G13D mutation. ICE CREAM: The Irinotecan Cetuximab Evaluation and the Cetuximab Response Evaluation Among Patients?	16/11/2012		No	No eligible patients were identified in clinics during the 70-day period
12/LO/1289	Pre-Exposure Option for reducing HIV in the UK: an open -label randomisation to immediate or deferred daily Truvada for HIV negative gay men	12/02/2013	14/03/2013	Yes	First patient recruited within 70 day benchmark
12/LO/0486	An Ascending Single Dose Study to Evaluate the Safety, Tolerability and Pharmacokinetics/Pharmacodynamics of PF-05280602: A Recombinant Factor Villa Variant (813), in adult Hemophilla A and B Subjects with or without inhibitors	21/11/2012		No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process
12/SC/0538	A Double-Blind, Placebo-Controlled, Randomized, Parallel Group, 12-Month Safety and Efficacy Trial of Leuco-methylthioninium bis(hydromethanesulfonate) in Subjects with Behavioral Variant Frontotemporal Dementia (bvFTD)	28/11/2012		No	Delays during NHS permissions and approvals process
13/LO/0696	Glycaemic index testing of carbohydrate containing foods using a standardised method	23/05/2013	28/05/2013	Yes	First patient recruited within 70 day benchmark
13/LO/0552	SinuSys Patency of Maxillary Sinus Ostia Study	05/06/2013	12/06/2013	Yes	First patient recruited within 70 day benchmark
13/LO/0126	Proportion of excision volume and length after treatment for cervical intra-epithelial lesions: cervical healing, quality of regenerated tissue, immunology and pregnancy outcome	23/04/2013	07/05/2013	Yes	First patient recruited within 70 day benchmark
13/LO/0425	A Randomized, Double-Blind, Controlled Study to Evaluate the Efficacy and Safety of the Combination of ABT-450/Ritonavir/ABT-267 (ABT-450/r/ABT-26)and BT-333 With and Without Ribavirin (RBV) in Treatment-Naive Adults with Genotype 1a Chronic?	18/04/2013	03/05/2013	Yes	First patient recruited within 70 day benchmark
12/LO/1278	Mechanisms of interplay between allergy and viruses in asthma: A human model of rhinovirus induced acute asthma exacerbations	23/01/2013	18/02/2013	Yes	First patient recruited within 70 day benchmark
12/LO/1570	Neuroinflammation in HIV individuals stable on cART: a [11C] PBR28 PET CT study	25/02/2013	28/03/2013	Yes	First patient recruited within 70 day benchmark
12/EE/0499	A Phase II Randomized, Dose Ranging, Clinical Trial to Evaluate the Safety, Tolerability, and Efficacy of Different Doses of MK-5172 When Administered Concomitantly with Peginterferon alfa-2b and Ribavirin in Treatment Na?ve Subjects with Chronic?	10/01/2013	18/02/2013	Yes	First patient recruited within 70 day benchmark
13/LO/0455	Retinal Embolisation in Transcatheter Aortic Valve Implantation: Pilot Study	17/05/2013	02/07/2013	Yes	First patient recruited within 70 day benchmark
12/LO/0331	ACE-EPIC: Enhancing the effects of pulmonary rehabilitation in COPD	29/10/2012	19/12/2012	Yes	First patient recruited within 70 day benchmark
11/SW/0036	Randomised Controlled Trial of the Efficacy and Mechanism of Levothyroxine Treatment on Pregnancy and Neonatal Outcomes in Women with Thyroid Antibodies	07/01/2013	04/03/2013	Yes	First patient recruited within 70 day benchmark
12/SC/0025	A 24-month, phase IIIb, openlabel, randomized, active controlled, 3-arm, multi-center study assessing the efficacy and safety of an individualized, stabilization criteria driven PRN dosing regimen with 0.5mg ranibizumab intravitreal injections...	21/08/2012	16/10/2012	Yes	First patient recruited within 70 day benchmark
12/LO/1320	Aneurysm Treatment using the HeliFX Aortic Securement System Global Registry	24/09/2012	28/11/2012	Yes	First patient recruited within 70 day benchmark

13/NW/0171	A Phase 2b, Randomized, Double-blind, Placebo-controlled Study Investigating the Efficacy and Safety of Inhaled CVT-301 (Levodopa Inhalation Powder) in Parkinson?s Disease Patients With Motor Response Fluctuations (OFF Phenomena)	13/05/2013	19/07/2013	Yes	First patient recruited within 70 day benchmark
12/LO/1748	A Phase 3, Multicenter, Randomized, Open-Label Study to Investigate the Efficacy and Safety of Sofosbuvir/GS-5885 Fixed Dose Combination + Ribavirin for 12 and 24 weeks in Treatment-Na?ve Subjects with Chroinc Genotype 1 HCV Infection	25/01/2013	04/04/2013	Yes	First patient recruited within 70 day benchmark
13/WM/0027	Venous Insufficiency and Neuromuscular Stimulation	15/04/2013	04/07/2013	No	No patients were consented during the 70-day period
12/LO/1174	AKTRES study: A Biologic Study of the early effects and determinants of AKT inhibition using GSK2110183 alongside chemotherapy in patients with Platinum RESistant Adenocarcinoma of the ovary	15/11/2012	04/02/2013	No	No patients were consented during the 70-day period
12/SC/0024	A 24 month, phase IIIb, open label, single arm, multi-center study assessing the efficacy and safety of an individualized, stabilization criteria driven PRN dosing regimen with 0.5mg ranibizumab intravitreal injections applied as monotherapy in...	19/06/2012	07/11/2012	No	No patients were consented during the 70-day period
13/WA/0064	A performance evaluation study of Urosens mcm5 Elisa test in the diagnosis of bladder cancer in patients presenting with haematuria and lower urinary tract symptoms with suspicion of malignancy	28/03/2013	02/07/2013	No	No patients were consented during the 70-day period / delays during NHS permissions and approvals process
13/LO/0115	A phase I clinical trial investigating immunisation strategies using DNA, MVA and rgp140 adjuvanted with GLAAF to maximise antibody responses	22/03/2013	28/06/2013	No	No patients were consented during the 70-day period / delays during NHS permissions and approvals process
12/LO/1361	A two-centre randomised controlled trial investigating the effect of remote ischaemic preconditioning (RIPC) on blood and myocardial biomarkers of stress and injury-related signalling in patients having isolated coronary artery bypass grafting?	07/11/2012	25/02/2013	No	No patients were consented during the 70-day period
12/SC/0014	Vasopressin vs. Noradrenaline as Initial therapy in Septic Shock	13/08/2012	09/02/2013	No	No patients were consented during the 70-day period
11/LO/0935	Genetic analysis for personalised medicine for morbid obesity	09/11/2011	02/08/2012	No	Delays during NHS permissions and approvals process
11/NW/0782	A phase III randomised trial of adjuvant chemotherapy versus observation in transitional cell carcinoma of the upper tract	21/05/2012	11/07/2013	No	No patients were consented during the 70-day period / delays during NHS permissions and approvals process
12/LO/1861	A Randomised, Open-label, Active-controlled, Multi-centre Study to Evaluate the Safety of Rivaroxaban and Vitamin K Antagonists in Subjects Undergoing Catheter Ablation for Atrial Fibrillation	10/12/2012		No	No eligible patients were identified in clinics during the 70-day period
12/SS/0007	A randomised Phase 2 Placebo-Controlled Study of LY2495655 in Patients with Advanced or Metastic Pancreatic Cancer Receiving Chemotherapy	10/12/2012		No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process
12/LO/0268	Open-Label, Phase 3b Study to Determine Efficacy and Safety of Telaprevir, Pegylated-Interferon-alfa-2a and Ribavirin in Hepatitis C Virus Treatment-Na?ve and Treatment Experienced Subjects with Genotype 1 Chronic Hepatitis C and Human Immuno?	17/12/2012		First patient recruited - awaiting confirmation of date	
12/LO/1769	Exploring the Physiological role of proportionate in glucose homoeostasis in man	15/01/2013		No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process
11/LO/0185	PROMIS Prostate MRI Imaging Study (MRC PR11) - Evaluation of Multi-Parametric Magnetic Resonance Imaging in the Diagnosis and Characterisation of Prostate Cancer	01/02/2013		No	Delays during NHS permissions and approvals process

12/LO/1156	A Phase III , Randomised, Double-blind , Controlled Study of Cabozantinib (XL184) vs Prednisone in Metastatic Castration-resistant Prostate Cancer Patients who have Received Prior Docetaxel and Prior Abiraterone or MDV3100?	06/02/2013		No	No eligible patients were identified in clinics during the 70-day period
12/LO/0777	Maraviroc Switch Cental Nervous System Study (March CNS) Substudy - a substudy of MARCH	12/02/2013		No	No eligible patients were identified in clinics during the 70-day period
09/H0707/88	Amino acid regimen and intravenous lipid composition in preterm parenteral nutrition: a randomised double blind controlled trial of Nutritional Evaluation and Optimisation in Neonates	12/02/2013		First patient recruited - awaiting confirmation of date	
10/H0706/65	A phase III trial comparing standard versus novel CRT as pre-operative treatment for MRI defined locally advanced rectal cancer (ARISTOTLE)	15/02/2013		No	No eligible patients were identified in clinics during the 70-day period
10/H0715/48	A Randomised Multicentre Accelerated Radiotherapy Study of Dose Escalated Intensity Modulated Radiotherapy vs Standard Dose Intensity Modulated Radiotherapy in Patients Receiving Treatment for Locally Advanced Laryngeal and Hypopharyngeal Cancers	15/02/2013		No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process
13/LO/0063	The acute effects of dietry components on central blood pressure	18/02/2013		First patient recruited - awaiting confirmation of date	
09/H0722/76	A randomised phase III trial of single fraction radiotherapy compared to multifraction radiotherapy in patients with metastatic spinal cord compression (SCORAD III)	26/02/2013		No	No eligible patients were identified in clinics during the 70-day period
12/LO/1896	Effect of Nike FuelBand on Exercise and Function in Claudicants	26/02/2013		No	No eligible patients were identified in clinics during the 70-day period
13/LO/0036	The effect of a low-carbohydrate, ketogenic diet verses a low-fat diet on weight loss and appetite regulation in type 2 diabetics	27/02/2013		No	No eligible patients were identified in clinics during the 70-day period
12/NW/0555	Randomized phase II study of BEZ235 or everolimus in advanced pancreatic neuroendocrine tumors	05/03/2013		No	No eligible patients were identified in clinics during the 70-day period
11/WS/0090	A Randomised Phase II study investigating pazopanib vs weekly paclitaxel in relapsed or progressive Transitional Cell Carcinoma (TCC) of the urothelium	08/03/2013		No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process
13/EE/0010	A phase I clinical trial to assess the safety and immunogenicity of three HIV GTU MultiHIV DNA immunisations administered via the Intramuscular, Intradermal and Transcutaneous routes in healthy male and female volunteers - CUTHIVAC 001?	08/03/2013		No	No eligible patients were identified in clinics during the 70-day period / delays during NHS permissions and approvals process
12/LO/1358	Randomized, Double-Blind, Phase 3 Study of TAS-102 Plus Best Supposrtive Care (BSC) Versus Placebo Plus BSC In Patients With Metastatic Colorectal Cancer Refractory To Standard Chemotherapies	11/03/2013		No	No eligible patients were identified in clinics during the 70-day period
12/NE/0342	A Study to Evaluate the Efficacy and Safety of GFT505 once daily on Steatohepatitis in Patients with Non-Alcoholic Steatohepatitis (NASH) - A Multicentre, Randomized, Double Blind, PlaceboControlled study, with an adaptive design to allow for?	11/03/2013		No	No eligible patients were identified in clinics during the 70-day period
12/NW/0682	A Double-blind, Randomized, Placebo-controlled, Multicenter Study to Evaluate Safety, Tolerability and Efficacy of AMG 145 on LDL-C in Subjects with Heterozygous Familial Hypercholesterolemia	13/03/2013		No	No eligible patients were identified in clinics during the 70-day period

13/LO/0006	A Phase 3, Open-label Study to Investigate the Efficacy and Safety of Sofosbuvir plus Ribavirin in Chronic Genotype 1, 2, 3 and 4 Hepatitis C Virus (HCV) and Human Immunodeficiency Virus (HIV) Co-infected Subjects	09/04/2013		First patient recruited - awaiting confirmation of date	
09/H0801/96	A Phase I Study Of Everolimus Therapy Before Nephrectomy In Metastatic Renal Cell Cancer (E-PREDICT)	24/04/2013		Still within 70 day period	
11/YH/0260	Combination FC plus Ofatumumab at Standard or Mega dose CLL	25/04/2013		Still within 70 day period	
13/LO/0223	Protexo Registry	25/04/2013		First patient recruited - awaiting confirmation of date	
12/LO/1981	A Randomized, Multicenter, Open-Label Phase III Study To Evaluate The Efficacy And Safety Of Trastuzumab Emtansine Versus Trastuzumab As Adjuvant Therapy For Patients With Her2-Positive Primary Breast Cancer Who Have Residual Tumor Present?	02/05/2013		First patient recruited - awaiting confirmation of date	
13/SC/0146	A phase III, randomised, open label, active controlled study of an interferon-free regimen of BI207127 in combination with Faldaprevir and Ribavirin compared to Telaprevir in combination with pegylated interferon and ribavirin in treatment na?ve?	03/05/2013		First patient recruited - awaiting confirmation of date	
12/LO/1823	A Randomized, Double-blind Phase 3B Study to Evaluate the Safety and Efficacy of Elvitegravir/Cobicistat/Emtricitabine/Tenofovir Disoproxil Fumarate Versus Ritonavir-Boosted Atazanavir Plus Emtricitabine/Tenofovir Disoproxil Fumarate in HIV 1?	17/05/2013		Still within 70 day period	
11/LO/2047	Outcome measures in sleeve gastrectomy after staple line reinforcement: Seamguard vs Duet TRS	22/05/2013		Still within 70 day period	
11/LO/1023	A 12-Month, Multicentre, Randomised, Parallel Group Study to Compare the Efficacy and Safety of Ozurdex Versus Lucentis in Patients with Branch Retinal Vein Occlusion	03/06/2013		Still within 70 day period	
12/NW/0711	Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, 12-Month Trial of Leuco-methylthioninium bis(hydromethanesulfonate) in Subjects with Mild to Moderate Alzheimer's Disease	06/06/2013		Still within 70 day period	
13/LO/0410	Preventing enduring behavioural problems in young children through early psychological intervention: A pilot study (Healthy Start Happy Start: Helping with Children's Behaviour)	07/06/2013		Still within 70 day period	
13/LO/0493	A randomised controlled trial of internet based cognitive behavioural therapy (CBT) versus treatment as usual (TAU) for pregnant women with symptoms of depression	10/06/2013		Still within 70 day period	