Id	Research Ethics Committee Reference Number	Integrated Research Application System Number	Name of Trial	First Patient Recruited?	Date of First Patient Recruited	Duration between Date Site Selected and Date Site Confirmed	Duration between Date Site Confirmed and First Patient Recruited	Duration between Date Site Selected and First Patient Recruited	Benchmark Met	Date Site Invited	Date Site Selected	HRA Approval Date	Date Site Confirmed By Sponsor	Date Site Confirmed	Date Site Ready To Start	Reasons for Delay	Comments	Reasons for delay correspond to:
108844	16/SW/0201	205118	A multi-centre, single arm, interventional Phase 4 study to evaluate a Treat and Extend regimen of intravitreal aflibercept for treatment of macular oedema secondary to central retinal vein occlusion	No		64			No	02/08/2016	19/10/2016	19/10/2016	14/12/2016	22/12/2016	11/01/2017	F - No patients seen	The DSS to DSC was over 40 days, this was due to capacity issues in the clinical setting	NHS Provider
108845	16/SC/0387	202827	Assessment of the effect of Positive Airway Pressure on energy and vitality in mild Obstructive Sleep Apnoea patients. The Merge Study	No		106			No	09/06/2016	23/11/2016	26/10/2016	31/01/2017	09/03/2017	09/03/2017	E - Staff availability issues		NHS Provider
108846	16/NW/0465	189554	Improving quality of life through the routine use of the Patient Concerns Inventory for head and neck cancer patients	Yes	08/02/2017	0	23	23	Yes	11/04/2016	16/01/2017	13/09/2016	16/01/2017	16/01/2017	16/01/2017			Please Select...
108847	16/WM/0006	192580	WHIST - Wound Healing In Surgery for Trauma. A Randomised Controlled Trial of standard wound management versus negative pressure wound therapy in the treatment of adult patients having surgical incisions for major trauma to the lower limb	Yes	03/02/2017	6	18	24	Yes	22/09/2016	10/01/2017	26/08/2016	30/01/2017	16/01/2017			This study was given HRA Approval for AUH to be added as a site. However, the original application was pre HRA Approval Process.	Please Select...
108848	16/NW/0786	186990	A feasibility study on integrating home-based metacognitive therapy for anxiety and depression in the cardiac rehabilitation pathway (PATHWAY STUDY 3)	Yes	10/04/2017	14	42	56	Yes	30/11/2016	13/02/2017	13/02/2017	13/02/2017	27/02/2017				Please Select...
108849	13/SC/0530	123453	An international multicentre randomised controlled trial of open versus laparoscopic liver surgery (hemihepatectomy and postero-superior segment resections).	Yes	14/09/2017	12	87	99	No	15/02/2017	07/06/2017	31/05/2017	14/06/2017	19/06/2017	19/06/2017	F - No patients seen	still within benchmark	Neither
108850	16/NS/0053	201313	A randomised controlled trial comparing laparoscopic cholecystectomy with observational/conservative management for preventing recurrent symptoms and complications in adults with uncomplicated symptomatic gall stones	No		33			No	28/02/2017	05/05/2017	01/09/2016	02/06/2017	07/06/2017		F - No patients seen	still within benchmark	Neither
108851	16/NS/0094	206213	Biolmpedance Spectroscopy to Maintain Renal Output: The BISTRO trial	Yes	26/06/2017	64	34	98	No	03/03/2017	20/03/2017	04/10/2016	15/05/2017	23/05/2017	19/06/2017	D - Sponsor Delays	Sponsor did not give Green Light until 19/06/2017	Sponsor
108852	16/EE/0065	192416	Survival improvement with Cholecalciferol in patients on dialysis -The SIMPLIFIED Registry trial	No		62			No	10/03/2017	10/03/2017	29/07/2016	10/03/2017	11/05/2017	26/06/2017	J - Other	Delays due to discussions with Sponsor regarding Excess Treatment Costs	Both
108853	13/LO/0968	98339	A randomized, double blind controlled trial comparing rituximab against intravenous cyclophosphamide in connective tissue disease associated interstitial lung disease	Yes	09/06/2017	9	35	44	Yes	18/12/2014	26/04/2017	10/08/2016	26/04/2017	05/05/2017	05/05/2017			Please Select...
108854	16/SC/0089	196789	A pragmatic randomised controlled trial to determine whether VV-ECCo2R in mechanically ventilated patients	Yes	26/06/2017	41	0	41	Yes	18/04/2016	16/05/2017	20/06/2016	20/06/2017	26/06/2017	26/06/2017		Still within benchmark	Please Select...

			with hypoxemic respiratory failure improves 90 day mortality															
108855	16/NW/0860	216349	Assessing and optimising the performance of low cost devices for measuring intravenous fluid delivery and response	Yes	17/05/2017	1	21	22	Yes	13/10/2016	25/04/2017	02/02/2017	25/04/2017	26/04/2017	26/04/2017			Please Select...
108906	15/WM/0268	180158	Randomised, open label study of rituximab/ibrutinib vs rituximab/chemotherapy in older patients with untreated mantle cell lymphoma	No		15			Within 70 Days	17/03/2017	14/08/2017	23/08/2016	24/08/2017	29/08/2017	31/08/2017	I - Rare diseases	There were pharmacy issues regarding excess treatment costs. This was relevant to the normal care pathway of the patients	NHS Provider
108907	16/NW/0496	169304	RESILIENT: RandomisEd, controlled, double blind Study to assess mechanistic effects of combination therapy of dapagliflozin with Exenatide LAR versus dapagliflozin alone in obese (BMI>30 kg/m2) patients with Type 2 diabetes mellitus.	No		29			Within 70 Days	27/10/2015	30/08/2017	12/06/2017	26/09/2017	28/09/2017		D - Sponsor Delays	Confirm CC from the Trust but green light still awaited from Sponsor	Sponsor
108908	16/NW/0517	188554	A phase III study to determine the role of ixazomib as an Augmented Conditioning therapy in salvage autologous stem cell transplant (ASCT) and as a post-ASCT Consolidation and maintenance strategy in patients with Relapsed multiple myeloma.	No		139			No	29/07/2016	15/02/2017	27/10/2016	17/07/2017	04/07/2017	04/07/2017	J - Other	This trial requires referral of patients to the Tranplant Centre. The Royal Liverpool is the Referral Centre. Our site could not open until the referral centre was open. This was the delay as the Royal did not open in line with our Trust.	Sponsor
108909	17/LO/0447	211176	A Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Study Assessing the Effect of Gelesis200 on Body Weight in Overweight and Obese Subjects with Prediabetes and Metformin-Treated Type 2 Diabetes (Protocol Num: GS-200-002)	Yes	30/08/2017	19	43	62	Yes	26/02/2017	29/06/2017	28/06/2017	06/07/2017	18/07/2017	01/08/2017			Please Select...
108910	17/NE/0096	222956	A randomised, double blind, placebo controlled multicentre Phase 2 dose ranging study to assess the safety and efficacy of multiple VAY736 doses administered subcutaneously in patients with moderate to severe primary Sjögren's syndrome	Yes	22/08/2017	39	22	61	Yes	29/03/2017	22/06/2017	26/05/2016	25/07/2017	31/07/2017	16/08/2017			Please Select...