

Research Ethics Committee Reference Number	Integrated Research Application System (IRAS) number	Submission Type	Name of Trial	Date of Receipt of Valid Research Application (old metric)	Date Site Invited (new metric)	Date Site Selected (new metric)	HRA Approval Date (new metric)	Date Site Confirmed By Sponsor (new metric)	Date Site Confirmed (new metric)	Date Site Ready to Start (new metric)	Date of First Patient Recruited	Non-Confirmation Status (new metric)	Duration between VRA / Date site selected and First Patient (days)	Benchmark Met	A - Permissions delayed/ denied	B - Suspended by sponsor	C - Closed by sponsor	D - Sponsor Delays	E - Staff availability issues	F - No patients seen	G - No patients consented	H - Contracting delays	I - Rare diseases	J - Other	Comments	Reasons for delay correspond to:	
13/WA/020127379		HRA Light	A Trial for Older Patients with Acute Myeloid Leukaemia and High Risk Myelodysplastic Syndrome		05/09/2016	21/11/2016	15/06/2016	11/11/2016	21/11/2016	13/12/2016	06/01/2017		46	Yes													
14/WA/1051154468		HRA Light	Adults with acute myeloid leukaemia or high-risk myelodysplastic syndrome (AML19)		02/09/2016	21/11/2016	15/06/2016	23/11/2016	29/11/2016	01/12/2016	05/12/2016		14	Yes													
15/LO/1302169660		HRA Light	OCTOPUS: Ovarian Cancer Trials of Weekly Paclitaxel - Umbrella StudyA Randomised, Phase II Umbrella Trial of a Weekly Paclitaxel +/- Novel Agents in Platinum-Resistant Ovarian Cancer		11/11/2016	09/01/2017	07/06/2016	03/05/2017	17/05/2017	07/06/2017	12/07/2017		184	No				Y							Internal capacity issues.	NHS Provider	
15/EE/0421191851		HRA Light	Pomalidomide in relapsed and refractory multiple myeloma		13/03/2017	30/03/2017	18/08/2016	24/04/2017	04/05/2017	08/06/2017	30/08/2017		153	No				Y	Y						Delays receiving study drug and lab kits at site.	Both	
BIO/MIL/01,135437		HRA Light	MILES - UK: Post Marketing, Multicenter, Single Arm, Observational Clinical Registry to Evaluate Safety and Efficacy of BioMime Sirolimus Eluting Stent System In All Comers Real World Population With Coronary Artery Stenosis in United Kingdom.		22/02/2017	10/05/2017	28/07/2016																				Within timeframe
15/NW/041167060		HRA Light	A randomised phase II trial of Cyclophosphamide and Dexamethasone in combination with Ixazomib, in relapsed or refractory multiple myeloma (RRMM) patients who have relapsed after treatment with thalidomide, lenolidomide and bortezomib.		07/03/2017	06/04/2017		19/05/2017	07/06/2017	28/06/2017				No											Delay due to additional contract with Myeloma UK - caused by both NHS		

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14/YH/1199	153953	HRA Light	GALACTIC: GA101 (obinutuzumab) monoclonal Antibody as Consolidation Therapy In CLL		18/05/2016	06/07/2016	15/06/2016					Sponsor declined site confirmation		N/A			Y								Letter received from trial office 11/07/2016 advising that recruitment into GALACTIC is being stopped. This is due to recruitment nationally being significantly slower than anticipated and trial office no longer think it is achievable to meet the recruitment target.	Neither
16/WM/003	180476	HRA Light	Accuracy of a rapid intrapartum test for maternal group B streptococcal colonisation and its potential to reduce antibiotic usage in mothers with risk factors.		28/12/2016	12/05/2017	23/08/2016	23/06/2017	05/07/2017									Y							Sub-category - delay intrinsic in study design Nature of delay - modifications to clinical testing guidelines	Sponsor
15/NS/0113	188563	HRA Light	The clinical and cost effectiveness of surgical interventions for stones in the lower kidney: The PUrE RCT- Percutaneous Nephrolithotomy (PNL), Flexible Ureterorenoscopy (FURS) and Extracorporeal Lithotripsy (ESWL) for lower pole Kidney stones.		04/05/2017	04/05/2017	09/06/2017	07/06/2017	14/06/2017						Y										Delays due to non-England lead review of study amendment to add WSHT as a site; amendment for WSHT as a site submitted 12/06/17 by sponsor, passed on by NRS to HRA on 04/07/17 and HRA confirmation (no HRA assessment required - non-substantial amendment) received 12/07/17. Advice received from NRS permissions stated "cannot open the study at the site until HRA Approval for the amendment is in place".	Neither