		Name of Trial	Recruited?	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	Invited	Date Site Selected	Date	Date Site Confirmed By Sponsor	Date Site Confirmed	Non- Confirmatic n Status	Date Site Ready To Start	Reasons for Delay	Reasons for delay correspon d to:
14/WA/1118		A Phase II/III trial of risk-stratified, reduced intensity adjuvant treatment in patients undergoing transoral surgery for Human papillomavirus (HPV)-positive oropharyngeal cancer	No					29/07/2015	10/01/2018	3					D - Sponsor Delays	Sponsor
17/WS/0030	191416		Yes	17/07/2017	10) 45	5 55	05/01/2017	23/05/2017	7 12/04/2017	/ 18/05/2017	02/06/2017	7	07/06/2017		
17/EE/0097		A Phase I Open-Labelled Trial investigating immunisation strategies using Ad4, MVA and rpg140 combinations in order to maximise antibody responses	Yes	09/03/2018	6	177	7 183	16/01/2017	07/09/2017	7 07/09/2017	16/01/2017	13/09/2017	,	06/10/2017	D - Sponsor Delays	Sponsor
16/EE/0065		Survival Improvement with Colecalciferol in Patients on Dialysis The SIMPLIFIED Registry Trial	No		140)		26/06/2016	17/11/2017	29/07/2016	6 26/06/2016	06/04/2018	8		D - Sponsor Delays	Sponsor
47/10/0524	20204.6			24/00/2047				07/02/2017	40/05/004-		07/00/0047	05/06/2047		05/06/2017	H - Contracting delays	6
17/LO/0524		Catheter ABlation of LOw VOltage regions in the treatment of persistent Atrial Fibrillation (ABLOVO-AF study)	Yes	24/08/2017	18	8 80		8 07/03/2017					, 	05/06/2017	B - Suspended by sponsor	Sponsor
17/EE/0330		A Prospective, Non-randomized, Consecutive Series, Multicentre, Observational Study to Evaluate the Clinical Outcome of Ceramic- on-Ceramic Hip Resurfacing Arthroplasty Using the H1 hip joint ceramic non-porous non-cemented prosthesis	Yes	26/09/2017	8	8 1	L <u>S</u>	17/09/2017	17/09/2017	7 22/09/2017	25/09/2017	25/09/2017	,	25/09/2017		
17/LO/0797		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.	No					28/11/2016	27/04/2017	01/08/2017	17/01/2018				B - Suspended by sponsor	Sponsor
17/SC/0411		A PROSPECTIVE, RANDOMIZED, DOUBLE-BLIND, PLACEBO- CONTROLLED PHASE 3 STUDY OF VGX-3100X (DNA PLASMID VECTORS EXPRESSING HPV-16 E6/E7, HPV-18 E6/E7) DELIVERED intramuscularly FOLLOWED BY EP WITH CELLECTRA® 5PSP FOR THE TREATMENT OF HISTOLOGICALLY PROVEN CIN2 OR CIN3 ASSOCIATED WITH HPV-16 AND/OR HPV-18	No					01/06/2016	23/08/2017						A - Permissions delayed/denied	Neither
1C/NF/041F	217015	A Dhace 2e. Onen Label Study to Evolute the Sefety	No		25			10/12/2010	01/00/2017	1 20/02/2017	10/12/2010	00/07/2017	Chancer de	lined site con	D - Sponsor Delays B - Suspended by sponsor	Chancar
16/NE/0415		A Phase 2a, Open-Label Study to Evaluate the Safety, Pharmacokinetics and Efficacy of the Combination of AL-335 and Odalasvir (ODV), with or without Simeprevir (SMV), in Treatment- Naïve Subjects with Genotype 1, 2 or 3 Chronic Hepatitis C infection with or without compensated Child Pugh A Cirrhosis	NO		35			10/12/2010	01/06/2017	28/02/2017	10/12/2010	06/07/2017	sponsor dec	linea site con	ib - Suspended by sponsor	Sponsor
17/LO/0465		ADJUNCTIVE GP ABLATION IN REDO-PVI STUDY (ADD-GP): PAROXYSMAL ATRIAL ARRHYTHMIAS AFTER PULMONARY VEIN ISOLATION ARE DRIVEN BY ECTOPY-TRIGGERING LEFT ATRIAL GANGLIONATED PLEXI	Yes	21/07/2017	54	37	7 91	. 24/02/2017	21/04/2017	7 21/04/2017	07/06/2017	14/06/2017			A - Permissions delayed/denied	NHS Provid
															F - No patients seen	
16/LO/1979		A randomised trial of non-Selective versus selective adjuvant Therapy in high risk Apparent sTage 1 Endometrial Cancer	Yes	19/03/2018	9	192	2 201	04/01/2017	30/08/2017	20/12/2016	06/09/2017	08/09/2017	/	08/09/2017	G - No patients consented	Neither
17/LO/0705		TRIAL ON RADICAL UPFRONT SURGERY IN ADVANCED OVARIAN CANCER AMENDMENT FOR GERMANY WITH EXTENDED PART FOR FRAGILITY AND LONG TERM QUALITY OF LIFE	Yes	04/09/2017	60) 34	1 94	02/06/2017	02/06/2017	7 28/07/2017	27/07/2017	01/08/2017		03/08/2017	G - No patients consented	Sponsor
															H - Contracting delays	
16/EM/0436		SINGLE ARM, STUDY OF ALXN1210 IN COMPLEMENT INHIBITOR TREATMENT-NAIVE ADULT AND ADOLESCENT PATIENTS WITH ATYPICAL HEMOLYTIC UREMIC SYNDROME (aHUS)	No		23			16/08/2017	16/08/2017	07/12/2016	16/08/2017	08/09/2017		15/09/2017	I - Rare diseases	Neither

47/10/4004		IN.	1	42				00/11/2017	1 10/10/2017	,	20/12/2017		D. Commentation	
17/LO/1991	172966 A comparative study of two popular weight-loss diets and their	No		42				08/11/201/	18/12/2017	'	20/12/2017		D - Sponsor Delays	Sponsor
	effect on the metabolic profiles of: i) Healthy overweight and													
	obese individuals ii) Overweight and obese breast cancer follow-													
16/WS/0197	up patients 186191 An adaptive multi-arm phase II trial of maintenance targeted	No		129			01/02/2017	20/05/2017	01/02/2017	02/08/2017	06/10/2017	12/10/2017	D - Sponsor Delays	Sponsor
10/ 003/0197	therapy after chemotherapy in metastatic urothelial cancer	NO		129			01/02/201/	50/03/201/	01/02/201/	02/06/201/	00/10/2017	12/10/2017	D - Spolisor Delays	Sponsor
													H - Contracting delays	
17/EE/0080	220727 A Randomized Phase 3 Study of AM0010 in Combination with	Yes	23/10/2017	29	96	125	30/03/2017	20/06/2017	09/06/2017	08/05/2017	19/07/2017	04/08/2017	I - Rare diseases	Neither
1772270000	FOLFOX Compared with FOLFOX Alone as Second-line Therapy in	105	23, 10, 2017	23	50	125	30/03/201/	20,00,201,	05,00,201,	00,00,201,	13/0//201/	04/00/201/		Nettrici
	Patients with Metastatic Pancreatic Cancer that has Progressed													
	During or Following a First-Line Gemcitabine Containing Regimen													
17/EE/0079	220827 A Randomized, Double-Blind, Placebo-Controlled, Phase 3 Study	Yes	23/01/2018	92	0	92	20/02/2017	23/10/2017	06/04/2017	20/02/2017	23/01/2018	23/01/2018	D - Sponsor Delays	Sponsor
17/22/00/5	to Evaluate the Safety and Efficacy of CCX168 (Avacopan) in	103	23/01/2010	52	Ű	52	20/02/201/	23/10/201/	00,04,201,	20/02/201/	23/01/2010	23/01/2010		5001301
	Patients with Anti-Neutrophil Cytoplasmic Antibody (ANCA)-													
	Associated Vasculitis Treated Concomitantly with Rituximab or													
	Cyclophosphamide/Azathioprine													
16/NS/0094		Yes	10/11/2017	65	30	05	22/02/2017	07/00/2017	04/10/2016	22/02/2017	11/10/2017	12/10/2017	D - Sponsor Delays	Chancar
10/103/0094	206213 BioImpedance Spectroscopy to Maintain Renal Output: The BISTRO Trial	res	10/11/2017	60	50	95	25/02/2017	07/08/2017	04/10/2016	25/02/2017	11/10/2017	12/10/2017	D - Sponsor Delays	Sponsor
17/55/0001		Vac	12/10/2017	62	99	101	20/02/2017		08/05/2017	20/06/2017	00/07/2017	00/07/2017	A - Permissions	Neither
17/EE/0081	219405 An open label, single arm pilot study of OncoSil™, administered to	res	13/10/2017	02	99	101	26/02/2017	05/05/201/	08/05/201/	29/06/201/	06/07/2017			Neither
	study participants with unresectable locally advanced pancreatic												delayed/denied	
	adenocarcinoma, given in combination with FOLFIRINOX or													
	gemcitabine+nab-paclitaxel chemotherapies.												L. Dava diasasas	
17/10/0124	210101 Account of a static latiffactor in action to with and stars hideau	Vee	15/00/2017	12	2		04/04/2010	02/05/2017	10/04/2017	04/04/2017	12/00/2017	12/05/2017	I - Rare diseases	
17/WM/0124	219181 Assessment of arterial stiffness in patients with end-stage kidney	Yes	15/06/2017	42	2	44	04/04/2016	02/05/201/	18/04/2017	04/04/201/	13/06/2017	13/06/2017		
17/10/0500	disease using shear-wave elastography			25			47/02/2017	44/04/2040	40/42/2047	47/02/2047	05 /02 /2010		C. No activate concentral	NI - tele - u
17/LO/0566	218514 An experimental medicine study to validate The 18 kD	No		25			1//03/201/	11/01/2018	19/12/201/	17/03/2017	05/02/2018		G - No patients consented	Neither
47/00/0400	Translocator Protein as a novel neuroimmunodulatory target							00/04/004-						-
17/SC/0130	33635 A Phase Ib Study Evaluating the Safety, Tolerability,	No						03/04/2017	ĺ			Site declined to participat	e	
	Pharmacokinetics, and Pharmacodynamics of GS-5829 in													
	Combination with Exemestane or Fulvestrant Followed by a													
	Parallel Randomized Phase II Study Comparing the Combination o	t												
	GS-5829 with Exemestane to Exemestane Alone and the													
	Combination of GS-5829 with Fulvestrant to Fulvestrant Alone in													
	Subjects with Advanced Estrogen Receptor Positive Breast Cancer													
17/NW/0175	222859 A Phase 1b Open-Label, Dose Escalation Study of PRTX-100 in	Yes	28/06/2017	58	6	64	20/10/2016	25/04/2017	19/05/2017	16/06/2017	22/06/2017	23/06/2017		
	Adult patients with Persistent/Chronic Immune													
	Thrombocytopenia													
16/EM/0465	214926 A Phase 2, Open-Label, Single-Agent, Multicenter Study to	No		285			14/03/2017	20/04/2017	19/01/2017	29/06/2017	30/01/2018		D - Sponsor Delays	Sponsor
	Evaluate the Efficacy and Safety of INCB054828 in Subjects With													
	Metastatic or Surgically Unresectable Urothelial Carcinoma													
	Harboring FGF/FGFR Alterations													
													H - Contracting delays	_
17/NW/0180	220257 A Phase 2, Open-Label, Single-Arm, Multicenter Study to Evaluate	No		108			15/03/2017	24/04/2017	04/04/2017	02/08/2017	10/08/2017	14/08/2017	D - Sponsor Delays	Sponsor
	the Efficacy and Safety of INCB054828 in Subjects With													
	Advanced/Metastatic or Surgically Unresectable													
	Cholangiocarcinoma Including FGFR2 Translocations Who Failed													
	Previous Therapy.													
17/LO/0746	218349 In comparison to usual care alone, does an adjunctive pre-surgery			1			20/02/2017	08/08/2017	01/08/2017	08/08/2017	09/08/2017	09/08/2017	B - Suspended by sponsor	Neither
	dietetic service improve the post-surgical nutritional outcomes fo	r												
	patients with pancreatic cancer? A pilot study													
													F - No patients seen	
Τ													G - No patients consented	
17/EM/0116	222773 A Multicenter, Open-Label Study to Evaluate the Safety and	Yes	18/07/2017	0	47	47	27/03/2017	01/06/2017	22/03/2017	27/03/2016	01/06/2017	11/07/2017	D - Sponsor Delays	Sponsor
	Tolerability of Tozadenant as Adjunctive Therapy in Levodopa-													
l I														
	Treated Patients with Parkinson's Disease Experiencing End of													

17/LO/0862	195365 The effects of fruits and vegetables on vascular function in prehypertensive participants: a pilot study	Yes	13/09/2017	2	53	55	21/04/2017	20/07/2017	04/07/2017	21/04/2017	22/07/2017	04/09/2017		
17/NE/0036	208636 A prospective, multicentre, randomised, controlled study evaluating SIR-Spheres Y-90 resin microspheres preceding standard cisplatin-gemcitabine (CIS-GEM) chemotherapy versus CIS-GEM chemotherapy alone as first-line treatment of patients with unresectable intrahepatic cholangiocarcinoma (SIRCCA)	Yes	08/01/2018	245	35	280	03/04/2017	03/04/2017	30/05/2017	30/11/2017	04/12/2017	07/12/2017	D - Sponsor Delays	Sponsor
													F - No patients seen	
													H - Contracting delays	
													I - Rare diseases	
17/LO/0783	225826 A Multicenter, Parallel-Group, Randomized, Cross-Over Trial to Compare the Efficacy of LUMINITY [®] and SonoVue [®] in the Evaluation of Left Ventricular Endocardial Border Definition	No		40			06/04/2017	24/08/2017	24/08/2017	25/09/2017	03/10/2017	16/01/2018	J - Other	NHS Provid
17/WS/0120	225090 A multi-center, double-blind, placebo-controlled, Phase 4 study patients with pulmonary arterial hypertension to assess the effe of selexipag on daily life physical activity and patient's self- reported symptoms and their impacts		11/01/2018	183	27	210	04/05/2017	15/06/2017	24/07/2017	13/12/2017	15/12/2017	21/12/2017	E - Staff availability issues	Sponsor
													H - Contracting delays	
													I - Rare diseases	
17/SC/0291	226509 The DEFINE PCI study: Physiologic assessment of coronary stenosis following PCI.	Yes	13/02/2018	139	63	202	09/03/2017	26/07/2017	29/08/2017	04/12/2017	12/12/2017	21/12/2017	E - Staff availability issues	Sponsor
													J - Other	
17/EM/0241	223736 Safety and efficacy analysis of FRED/FRED Jr embolic device in aneurysm treatment	Yes	23/11/2017	62	65	127	05/07/2017	19/07/2017	08/08/2017	18/09/2017	19/09/2017	19/09/2017	F - No patients seen	NHS Provid
													H - Contracting delays	
16/EE/0463	214371 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 pre/postmenopausal women with hormone receptor-positive (HR+) HER2-negative (HER2-) advanced breast cancer (aBC) with no prior hormonal therapy for advanced disease.		20/10/2017	71	28	99	12/06/2017	13/07/2017	30/01/2017	18/09/2017	22/09/2017		A - Permissions delayed/denied	Both
													H - Contracting delays	
17/SC/0253	226685 A Randomized, Open-Label, Phase 3 Study of Abemaciclib Combined with Standard Adju-vant Endocrine Therapy versus Standard Adjuvant Endocrine Therapy Alone in Patients with Hig Risk, Node Positive, Early Stage, Hormone Receptor Positive, Human Epidermal Receptor 2 Negative, Breast Cancer.	No h		187			30/05/2017	31/05/2017	05/07/2017	30/11/2017	04/12/2017	03/01/2018	D - Sponsor Delays	Sponsor
													G - No patients consented	
17/LO/1169	224716 Efficacy and usability of Ostom-i device in patients with ileostom A pilot study	iy. No		1			25/09/2017	22/11/2017	20/10/2017		23/11/2017		G - No patients consented	Neither
17/NW/0254	225698 A Phase III, Randomized, Double-Blind, Placebo-Controlled Clinic Trial of Pembrolizumab (MK-3475) as Monotherapy in the Adjuvant Treatment of Renal Cell Carcinoma Post Nephrectomy	al Yes	27/11/2017	85	46	131	05/05/2017	19/07/2017	17/07/2017	25/09/2017	12/10/2017		A - Permissions delayed/denied	Sponsor
													D - Sponsor Delays	
													F - No patients seen	
													H - Contracting delays	
17/NE/0149	224550 (M16-298) randomized, double-blind, placebo-controlled phase study of rovalpituzumab tesirine (Rova-T) as maintenance therap following four cycles of firstline therapy with a platinum doublet in subjects with extensive stage disease (ED) SCLC	ру		142			10/05/2017	20/06/2017	19/06/2017	06/09/2017	09/11/2017		A - Permissions delayed/denied	Both
													H - Contracting delays	
17/LO/0939	219566 Patient Empowerment Through Predictive Personalised Decision Support (PEPPER)	Yes	09/01/2018	35	22	57	12/09/2017	13/11/2017	08/11/2017	12/09/2017	18/12/2017	18/12/2017		

16/WM/0448	212745 A Phase 3, Multicenter, Randomized, Open-label, Active- controlled Study to Assess the Efficacy and Safety of Maribavir	No		222			05/04/2017	05/04/2017	08/02/2017	23/10/2017	13/11/2017	16/11/2017	D - Sponsor Delays	Sponsor
	Treatment Compared to Investigator-assigned Treatment in Transplant Recipients with Cytomegalovirus (CMV) Infections that													
	are Refractory or Resistant to Treatment with Ganciclovir,													
	Valganciclovir, Foscarnet, or Cidofovir.	Vac 10	/01/2019	101		107	07/00/2017	00/07/2017	05/07/2017		14/11/2017	17/11/2017	D. Changer Delays	Chancer
17/ES/0051	223060 CANC 33870: A Phase II Open-Label Study of NUC-1031 in Patients with Platinum-Resistant Ovarian Cancer	Yes 19	/01/2018	131	66	197	07/06/2017	06/07/2017	05/07/2017		14/11/2017	1//11/201/	D - Sponsor Delays	Sponsor
													H - Contracting delays	
17/LO/1095	225278 Investigation of the metabolic effects of duodenal resurfacing on	No		39			09/06/2017	09/02/2018	04/09/2017	09/06/2017	20/03/2018	21/03/2018		
17/EE/0264	insulin resistant women with polycystic ovarian syndrome	No		75			02/00/2017	12/00/2017	12/09/2017	20/11/2017	27/11/2017	27/11/2017	D - Sponsor Delays	Chancer
17/66/0204	228153 A Phase III, Randomized, Multi-Center, Open-Label, Comparative Global Study to Determine the Efficacy of Durvalumab or	NO		75			02/06/2017	13/09/201/	12/09/201/	20/11/201/	27/11/2017	2//11/201/	D - Sponsor Delays	Sponsor
	Durvalumab and Tremelimumab in Combination With Platinum-													
	Based Chemotherapy for First-Line Treatment in Patients With													
	Metastatic Non-Small-Cell Lung Cancer (NSCLC) (POSEIDON)													
													F - No patients seen	
													H - Contracting delays	
17/EE/0177	220722 A Phase 3 Randomized, Controlled, Open-label Study of Selinexor,	Yes 03	/10/2017	63	7	70	31/05/2017	25/07/2017	24/07/2017	18/09/2017	26/09/2017	27/09/2017		
1772270177	Bortezomib, and Dexamethasone (SVd) versus Bortezomib and Dexamethasone (Vd) in Patients with Relapsed or Refractory		,10,2017		,	70	51/05/2017	23/07/2017	24/07/2017	10/03/2017	20/03/2017	21/05/2017		
47/00/0554	Multiple Myeloma (RRMM).	N		100			24 /05 /2017	40/40/2047	1 22/02/2010	24/02/2010	101/2010		D. Courses Dalaus	C
17/SC/0554	225522 A Phase III, multicenter, randomized controlled study to compare	NO		188			31/05/2017	10/10/201/	23/02/2018	21/03/2018	16/04/2018		D - Sponsor Delays	Sponsor
	safety and efficacy of a haploidentical HSCT and adjunctive													
	treatment with ATIR101, a T-lymphocyte enriched leukocyte													
	preparation depleted ex vivo of host alloreactive T-cells, versus a													
	haploidentical HSCT with post-transplant cyclophosphamide in													
	patients with a hematologic malignancy (HATCY study).												H - Contracting delays	
													J - Other	
16/WA/0156	204506 Minimally invasive thoracoscopically-guided right	Yes 13	/03/2018	205	26	231	14/06/2017	25/07/2017	19/06/2016	31/01/2018	15/02/2018	15/02/2018	A - Permissions	Sponsor
10, WA 0130	minithoracotomy versus conventional sternotomy for mitral valve repair: a multicentre randomised controlled trial (UK Mini Mitral).	1 1	,03,2010	203	20	231	14/00/2017	23/07/2017	15/00/2010	51/01/2010	15/02/2010	13/02/2010	delayed/denied	5001301
													D - Sponsor Delays	
													E - Staff availability issues	
													H - Contracting delays	
17/LO/1590	229278 A Phase III, Randomized, Double-Blind, Clinical Trial of	No		121			11/07/2017	01/11/2017	27/10/2017		02/03/2018	02/03/2018	A - Permissions	Sponsor
	Pembrolizumab (MK-3475) plus Chemotherapy (XP or FP) versus												delayed/denied	
	Placebo plus Chemotherapy (XP or FP) as Neoadjuvant/Adjuvant													
	Treatment for Subjects with Gastric and Gastroesophageal													
	Junction (GEJ) Adenocarcinoma (KEYNOTE-585).													
													D - Sponsor Delays	
													H - Contracting delays	
17/SW/0127	225959 A multicentre randomised trial of First Line treatment pathways	No	T	205			19/06/2017	19/06/2017	03/07/2017	09/01/2018	10/01/2018		J - Other	NHS Provid
	for newly diagnosed Immune Thrombocytopenia: Standard													
	steroid treatment versus combined steroid and mycophenolate.													
17/LO/1441	219916 SYNCope EDUcation for patients during tilt-table tests	Yes 20	/03/2018	26	36				12/09/2017		12/02/2018			
17/NE/0174	224141 A PERFORMANCE EVALUATION STUDY OF ARQUER'S MCM5 ELISA	Yes 06	6/09/2017	65	5	70	27/06/2017	28/06/2017	13/06/2017	08/08/2017	01/09/2017	01/09/2017		
	TEST TO AID IN THE MONITORING OF BLADDER CANCER RECURRENCE													
17/LO/1457	2277777 Randomised controlled trial of mechanochemical ablation versus	Yes 06	6/11/2017	9	5	14	01/08/2017	23/10/2017	19/09/2017	30/10/2017	01/11/2017	30/10/2017		
, ,,	cyanoacrylate adhesive for the treatment of varicose veins		, ,	-	-		,,	, ,,=-=,		, ,,=-=,	, ,			
17/LO/1447	226100 The effectiveness of a focused point-of-care duplex ultrasound	No		60			30/08/2017	06/11/2017	21/12/2017	15/12/2017	05/01/2018		A - Permissions	Neither
, , ,	scan as a bedside screening tool to detect peripheral arterial						,,	, ,====	, ,====,	, ,====			delayed/denied	
	disease in diabetes.												, ,	

17/WS/0192		Randomised controlled trial of foam sclerotherapy versus ambulatory phlebectomy for the treatment of varicose vein tributaries	Yes	22/01/2018	38	61	99	21/08/2017	15/10/2017	15/10/2017		22/11/2017		E - Staff availability issues	Neither
												1		F - No patients seen	
														G - No patients consented	
														J - Other	
14/LO/0801		Improving radical treatment through MRI evaluation of pelvic sigmoid cancers	No		225			08/08/2017	08/08/2017	15/06/2016	21/03/2018	3 21/03/2018	27/03/2018	D - Sponsor Delays	Sponsor
														H - Contracting delays	
17/LO/1247		RANDOMIZED, OPEN LABEL, MULTICENTRE, PHASE 3 STUDY OF ROVALPITUZUMAB TESIRINE COMPARED WITH TOPOTECAN FOR SUBJECTS WITH ADVANCED OR METASTIC DLL3 SMALL CELL LUNG CANCER (SCLC) WHO HAVE FIRST DISEASE PROGRESSION DURING OR FOLLOWING FRONT LINE PLATINUM -BASED CHEMOTHERAPY	No		164			30/01/2017	22/08/2017	15/08/2017		02/02/2018		A - Permissions delayed/denied	NHS Provid
1 - 4 - 4 - 4 - 4				1 - / 1 / 2 0 / -								1 - 1 - 0 - 0 - 0		H - Contracting delays	
17/LO/1040		p53 Suppressor Activation in Platinum-Resistant High Grade Serous Ovarian Cancer, a Phase II Study of Systemic Pegylated Liposomal Doxorubicin Chemotherapy With APR-246	Yes	17/11/2017	1	31	32		16/10/2017	04/09/2017		17/10/2017	27/10/2017		
17/YH/0311		A Modular, Multipart, Multiarm, Open-label, Phase I/IIa Study to Evaluate the Safety and Tolerability of CT7001 Alone and in Combination with Anti-cancer Treatments in Patients with Advanced Malignancies	Yes	28/11/2017	79	4	83	29/08/2017	06/09/2017	30/10/2017		24/11/2017	24/11/2017	D - Sponsor Delays	Both
														H - Contracting delays	
14/LO/0843		The role of MRI in improving surgical technique and outcomes in exenterative pelvic surgery for locally extensive primary or recurrent rectal cancer	No		201			01/09/2017	01/09/2017	18/01/2017	21/03/2018	3 21/03/2018	27/03/2018	D - Sponsor Delays	Sponsor
														H - Contracting delays	
17/WS/0180		A Phase 111b, Randomised, Double-blind, Placebocontrolled, MulticentreStudy of Olaparib Maintenance Retreatment in Patients with EpithelialOvarian Cancer Previously Treated With a PARPi and Responding toRepeat Platinum Chemotherapy (OReO)	No		174			11/09/2017	13/10/2017	02/02/2018		05/04/2018	29/03/2018	H - Contracting delays	Sponsor
17/EE/0340		INTERIM: a randomised phase II feasibility study of INTERmittent versus continuous dosing of oral targeted combination therapy (BRAF+ MEKi) in patients with BRAFV600 mutant stage 3 unresectable or metastatic Melanoma.	No		82			01/08/2017	25/10/2017	15/10/2017	03/01/2018	3 15/01/2018		A - Permissions delayed/denied	NHS Provid
17/SC/0394	229032	An Open-Label, Randomized, Phase 2 Dose-Finding Study of Pacritinib in Patients with Thrombocytopenia and Primary Myelofibrosis, Post-Polycythemia Vera Myelofibrosis, or Post- Essential Thrombocythemia Myelofibrosis Previously Treated with Ruxolitinib.	Yes	19/12/2017	101	26	127	07/08/2017	14/08/2017	23/10/2017		23/11/2017	12/12/2017	D - Sponsor Delays	Sponsor
17/LO/1149	220763	A Phase 3, multicenter, randomized, double-blind, double- dummy, active-controlled study to assess the efficacy and safety of maribavir compared to valganciclovir for the treatment of cytomegalovirus (CMV) infection in hematopoietic stem cell transplant recipients.	No		194			26/07/2017	26/07/2017	12/09/2017	09/10/2017	05/02/2018	06/03/2018	D - Sponsor Delays	Sponsor
17/EM/0338	229242	A phase 3, multi-center, open-label, randomized study of oral ABL001 versus bosutinib in patients with Chronic Myelogenous Leukemia in chronic phase (CML-CP), previously treated with 2 or more tyrosine kinase inhibitors	No					04/09/2017	07/02/2018	24/01/2018				D - Sponsor Delays	Sponsor
17/LO/1320	229845		No		153			10/08/2017	10/08/2017	27/10/2017	22/12/2017	/ 10/01/2018		A - Permissions delayed/denied	Sponsor
			ļ								ļ	-		D - Sponsor Delays	_
														H - Contracting delays	
17/SC/0408		A Randomized, Double-Blind, Placebo-Controlled Phase 3 Study of Itacitinib or Placebo in Combination with Corticosteroids for the Treatment of First-Line Acute Graft-Versus-Host Disease.	No		200			18/09/2017	18/09/2017	19/09/2017	06/04/2018	8 06/04/2018		D - Sponsor Delays	Sponsor

17/LO/1429	223671 A Two-Part Phase 1/2 Study to Determine Safety, Tolerability,	No		191		06/09/2017	06/09/2017	29/01/2018	12/01/2018	16/03/2018	21/03/2018	D - Sponsor Delays	Sponsor
17/10/1429	Pharmacokinetics, and Activity of K0706, a Novel Tyrosine Kinase	NO		191		00/03/201/	00/03/201/	23/01/2010	12/01/2010	10/03/2018	21/03/2018	D - Spolisor Delays	5001501
	Inhibitor (TKI), in Healthy Subjects and in Subjects with Chronic												
	Myeloid Leukemia (CML) or Philadelphia Chromosome Positive												
	Acute Lymphoblastic Leukemia (Ph+ ALL).												
												J - Other	
17/LO/1847	216867 A randomised phase II trial of Selinexor, cyclophosphamide and	No				22/11/2017	14/02/2018	04/01/2018				D - Sponsor Delays	Sponsor
	prednisone vs cyclophosphamide and prednisone in relapsed or												
	refractory multiple myeloma (RRMM) patients.												
17/LO/1500	224823 A Phase 3, Multicenter, Double-Blind, Randomized, Placebo-	No				05/02/2018	13/02/2018					D - Sponsor Delays	Sponsor
, , .,	controlled Study of AG-120 in Combination with Azacitidine in					,-,-	-,-,-						
	Subjects = 18 Years of Age with Previously Untreated Acute												
	Myeloid Leukemia with an IDH1 Mutation Who are candidates for												
	Non-intensive Therapy.												
17/SC/0201	Tazemetostat Rollover Study (TRuST): An Open-Label, Rollover	No				29/01/2018	22/03/2018	18/10/2017					
17,00,0201	Study.					20,01,2010	22,00,2010	10, 10, 201,					
17/EM/0371	229496 Strategic Management to Optimize Response To Cardiac	No				10/07/2017	26/03/2018	21/11/2017					
	Resynchronization Therapy												
17/LO/1695	234684 A Phase 3 Open-Label, Single-Arm Study To Evaluate The Efficacy	No		65		09/10/2017	09/10/2017	24/11/2017		13/12/2017		A - Permissions	Neither
	and Safety of BMN 270, an Adeno-Associated Virus Vector-											delayed/denied	
	Medicated Gene Transfer of Human Factor VIII in Haemphilia A												
	Patients with Residual FVIII Levels 1 IU/dl Receiving Prophylactic												
	FVIII Infusions.												
16/EM/0382	210424 A multicenter, randomized, double-blind, active-controlled study	No		40		12/10/2017	23/02/2018	05/12/2016	08/02/2018	04/04/2018			
-, ,	to evaluate the effects of LCZ696 compared to valsartan on			-		, , -, -	-,-,-	, , , , , , , , , , , , , , , , , , , ,					
	cognitive function in patients with chronic heart failure and												
	preserved ejection fraction.												
17/LO/1918	229435 Does Neuromuscular Electrical Stimulation Improve the Absolute	Yes	08/03/2018	45	20	65 09/11/2017	02/01/2018	29/01/2018		16/02/2018	16/02/2018	E - Staff availability issues	
1,720,1010	Walking Distance in Patients with Intermittent Claudication		00/00/2010	15	20	05 05/11/201/	02,01,2010			10,02,2010	10,02,2010		
	(NESIC) compared to best available treatment? A Multicentre												
	Randomised Controlled Study												
17/LO/1592	225931 AN OPEN-LABEL, MULTICENTER EXTENSION STUDY IN PATIENTS	No		89		15/09/2017	20/09/2017	07/11/2017	12/12/2017	18/12/2017	10/12/2017	A - Permissions	Neither
1771071332	PREVIOUSLY ENROLLED IN A GENENTECHâ [*] AND/OR F.			05		15/05/2017	20,03,201,	0,,11,201,	12/12/2017	10/12/2017		delayed/denied	Neither
	HOFFMANN-LA ROCHE LTDâ [^] SPONSORED ATEZOLIZUMAB STUDY											delayed/deliled	
17/NE/0151	223815 Phase 3, Multicenter, Randomized, Open-Label Study of	No		135		01/11/2017	01/11/2017	10/07/2017	07/03/2018	16/03/2018	06/04/2018	D - Sponsor Delays	Sponsor
17/1010101	Guadecitabine (SGI-110) versus Treatment Choice in Adults with			133		01/11/2017	01/11/201/	15/0//201/	07/03/2010	10/03/2010	00/04/2018		5001301
	Previously Treated Acute Myeloid Leukemia (SGI-110-06).												
17/LO/1929	233885 A Phase 3 Randomized, Open-Label Clinical Study to Evaluate the	No		148		02/11/2017	12/11/2017	05/02/2018	27/02/2019	10/04/2018		H - Contracting delays	NHS Provi
17/10/1929		NO		140		02/11/2017	13/11/201/	03/02/2018	27/03/2018	10/04/2018		H - Contracting delays	
	Efficacy and Safety of Pembrolizumab plus Epacadostat,												
	Pembrolizumab, and the EXTREME Regimen as First line												
	Treatment for Recurrent or Metastatic Head and Neck Squamous												
17/55/0401	Cell Carcinoma (KEYNOTE-669/ECHO-304)	Na		74		00/11/2017	24/11/2017		00/11/2017	00/02/2010		A Dermineiere	Naithau
17/EE/0481	226412 Pre-Implantation trial of Histopathology In renal Allografts	No		74		08/11/2017	24/11/2017	05/01/2018	08/11/201/	06/02/2018		A - Permissions	Neither
18/10/0100	237129 A Study of Flat and Circadian Insulin Infusion Rates in Continuous	No		16		10/01/2019	27/02/2018			15/03/2018		delayed/denied	
18/LO/0196	Subcutaneous Insulin Infusion (CSII) in Adults with Type 1	NO		10		10/01/2018	27/02/2010			15/05/2018			
17/10/0119		No		7		02/11/2017	14/12/2017	28/04/2017	14/12/2017	21/12/2017		C No notionts concented	Naithar
17/LO/0118	220143 Maternal Moments: Investigating music listening of well-being in pregnancy	NO		/		03/11/2017	14/12/2017	28/04/2017	14/12/201/	21/12/2017		G - No patients consented	Neither
17/NW/0351	224954 A phase II single arm clinical trial of a Tailored ImmunoTherapy	No					12/09/2017	,				A - Permissions	NHS Provi
1,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	Approach with Nivolumab in subjects with metastatic or advanced						12,00,201,					delayed/denied	
	Renal Cell Carcinoma											ucity cu ucincu	
17/EE/0368		No				12/10/2017	28/03/2018	10/11/2017	-				
1,722,0300	(Beta Blockade)	10				12/10/2017	20/03/2010	10,11,201,					
17/EE/0480	235288 A Phase 3 Randomized, Double-Blind Trial of Pembrolizumab (MK-	No		117		27/11/2017	14/12/2017	05/02/2018	27/03/2018	10/04/2018		D - Sponsor Delays	Sponsor
1, 22, 0400	3475) in Combination with Epacadostat (INCB024360) or Placebo			±±/		2,,11,2017	,,,,	33, 02, 2010					5,01,501
	in Participants with Cisplatin-ineligible Urothelial Carcinoma												
	(KEYNOTE-672/ECHO-307)												
L		1							1				

														H - Contracting delays	
17/LO/2059		A Phase 3 Randomized, Double-Blind Clinical Study of Pembrolizumab + Epacadostat vs Pembrolizumab + Placebo as a Treatment for Recurrent or Progressive Metastatic Urothelial Carcinoma in Patients who have Failed a First-Line Platinum- containing Chemotherapy Regimen for Advanced/Metastatic Disease (KEYNOTE-698/ECHO-303)	No		103			27/11/2017	14/12/2017	29/01/2018	20/03/2018	27/03/2018		D - Sponsor Delays	Sponsor
														H - Contracting delays	
17/SC/0641	238111	A Phase 1/2 Safety, Tolerability, and Efficacy Study of BMN 270, an Adeno-Associated Virus Vector–Mediated Gene Transfer of Human Factor VIII in Hemophilia A Patients with Residual FVIII Levels = 1 IU/dL and Pre-existing Antibodies Against AAV5	No				(08/12/2017	08/12/2017				Sponsor declined site cor	•	
18/WM/0022	237623	AG-348 in Regularly Transfused Adult Subjects with PK Deficiency	No						19/01/2018					D - Sponsor Delays	Sponsor
18/LO/0054		Microneedle sensing of beta-lactam antibiotic concentrations in human interstitial fluid	No		14		(06/07/2017	21/03/2018	14/02/2018	06/07/2017	04/04/2018			
17/LO/0103		A Multicenter, Double-Blind, Double-Dummy Study to Explore the Long-Term Safety of TEV-48125 for the Prevention of Cluster Headache	No		15		:	20/06/2017	20/12/2017	05/07/2017		04/01/2018		G - No patients consented	Neither
17/LO/0089	219229	A Phase 2, Multicenter, Open-Label Study to Evaluate the Efficacy and Safety of Sofosbuvir/Velpatasvir for 12 Weeks in Subjects with Chronic HCV Infection Who are on Dialysis for End Stage Renal Disease	Yes	15/07/2017	21	29	50 2	27/01/2017	26/05/2017	27/03/2017	27/01/2017	16/06/2017	16/06/2017		
17/LO/0102		A Multicenter, Randomized, Double-Blind, Double-Dummy, Placebo-Controlled, Parallel-Group Study Comparing the Efficacy and Safety of 2 Dose Regimens (Intravenous/ Subcutaneous and Subcutaneous) of TEV-48125 versus Placebo for the Prevention of CHRONIC Cluster Headaches (CCH)	No		11		:	12/12/2016	22/12/2017	14/07/2017	12/12/2016	02/01/2018		D - Sponsor Delays	Sponsor
16/EE/0421		A Comparison of Bimatoprost SR to Selective Laser Trabeculoplasty in Patients with Open-Angle Glaucoma or Ocular Hypertension	No					15/02/2017	25/04/2017	13/01/2017				D - Sponsor Delays	Sponsor
														H - Contracting delays	
17/LO/0444		A Follow-up Study to Assess Resistance and Durability of Response to AbbVie Direct-ActingAntiviral Agent (DAA) Therapy (ABT-493 and/or ABT-530) in Subjects Who Participated in Phase 2 or 3 Clinical Studies for the Treatment of Chronic Hepatitis C Virus (HCV) Infection	Yes	19/06/2017	1	10	11 (08/03/2017	08/06/2017	05/05/2017	08/03/2017	09/06/2017	09/06/2017		
17/LO/0742		A randomised controlled comparison of effectiveness of facemask pre-oxygenation and Transanal Humidified Rapid-Insufflation Ventilatory Exchange (THRIVE) in bariatric patients undergoing general anesthesia	No		3			15/02/2017	10/07/2017	29/06/2017	13/07/2017	13/07/2017	17/07/2017	D - Sponsor Delays	Sponsor
16/WM/0285		A Double Blind, Randomized, Placebo Controlled, Multicenter Study to Evaluate Safety, Tolerability, and Efficacy on LDL-C of Evolocumab (AMG 145) in Subjects with HIV and with Hyperlipidemia and/or Mixed Dyslipidemia	Yes	01/08/2017	16	12	28 :	17/03/2017	04/07/2017	22/06/2017	17/03/2017	20/07/2017	05/07/2017		
17/LO/0401		An Open-Label, Multicenter Study to Evaluate the Efficacy and Safety of ABT-493/ABT-530 in Combination with Sofosbuvir and Ribavirin in Chronic Hepatitis C (HCV) Infected Subjects Who Have Experienced Virologic Failure in AbbVie HCV Clinical Studies.	Yes	06/06/2017	22	13	35 (02/05/2017	02/05/2017	23/03/2017	21/03/2017	24/05/2017	31/05/2017	,	
17/LO/0096	220205	A Phase 3b, 12-month, Open-label, Multicenter Study to Evaluate the Efficacy and Safety of BIIB019, Daclizumab, in Subjects with Relapsing-Remitting Multiple Sclerosis (RRMS) Switching from Natalizumab (SUSTAIN)	No		45		:	25/01/2017	11/04/2017	11/04/2017		26/05/2017	03/07/2017	B - Suspended by sponsor	Sponsor
17/LO/1003		Perforated Punctal Plug as an alternative to the three snip punctoplasty for the treatment of acquired lacrimal punctum stenosis	Yes	04/09/2017	0	24	24 2	22/03/2017	11/08/2017	01/08/2017	11/08/2017	11/08/2017	23/08/2017		
17/LO/1036	220972	Feasibility study for the use of intraoperative single dose radiotherapy for locally advanced and recurrent pelvic cancer	No		12		2	20/04/2017	05/10/2017	06/09/2017		17/10/2017	17/10/2017	I - Rare diseases	Neither

16/WM/0514			No					30/03/2017	06/04/2017	06/04/2017	30/03/2017		Sponsor declined site con	J - Other	Sponsor
17/LO/0095	216048	using ENdocuff Optimisation of Mucosal Abnormalities A tailored, cognitive behavioural approach intervention for mild to moderate anxiety and/or depression in people with chronic obstructive pulmonary disease (COPD): A randomised controlled trial.	No		108				10/04/2017	08/03/2017		27/07/2017		D - Sponsor Delays	Sponsor
														H - Contracting delays	
16/NW/0877		A Safety and Tolerability Study of LY3303560 in Alzheimer's Disease	Yes	06/10/2017	6	66	72	19/04/2017	26/07/2017	05/04/2017	19/04/2017	01/08/2017	01/09/2017	D - Sponsor Delays	Sponsor
16/SS/0137	199347	Alteplase-Tenecteplase Trial Evaluation for Stroke Thrombolysis 2	No		129			24/04/2017	07/11/2017	11/01/2017	24/04/2017	16/03/2018		D - Sponsor Delays	Sponsor
17/WM/0425		A phase III trial evaluating the efficacy and safety of the SQ house dust mite (HDM) sublingual immunotherapy (SLIT)-tablet in children and adolescents with HDM allergic asthma			55				24/11/2017				08/03/2018	A - Permissions delayed/denied	Neither
17/SC/0122		AR101 Trial in Europe Measuring Oral Immunotherapy Success in Peanut Allergic Children	Yes	18/07/2017	63	6	69		10/05/2017				14/07/2017	H - Contracting delays	
16/SC/0338		Prospective Interruption of Therapy towards a Cure for HIV (PITCH) Pilot Study	No		196			09/05/2017	09/08/2017	22/11/2016	17/10/2017	21/02/2018		H - Contracting delays	Sponsor
16/YH/0157		PLATO - PersonaLising Anal cancer radioTherapy dOse - Incorporating Anal Cancer Trials (ACT) ACT3, ACT4 and ACT5	No		128			26/08/2016	6 06/09/2017	20/07/2016		12/01/2018	18/01/2018	D - Sponsor Delays	Sponsor
														F - No patients seen	
17/LO/1129	226325	An adaptive orthosis for hand osteoarthritis; feasibility and prototype	Yes	04/10/2017	7	20	27	08/09/2017	07/09/2017	07/09/2017	08/09/2017	14/09/2017	14/09/2017		
17/NE/0165		AN OPEN-LABEL, SINGLE-ARM STUDY TO EVALUATE THE EFFECTIVENESS AND SAFETY OF OCRELIZUMAB IN PATIENTS WITH EARLY STAGE RELAPSING REMITTING MULTIPLE SCLEROSIS	Yes	21/11/2017	133	21	154	22/05/2017	20/06/2017	07/09/2017	22/05/2017	31/10/2017	14/11/2017	D - Sponsor Delays	Sponsor
														H - Contracting delays	
16/SW/0120	201715	Study of the optimum duration of acoustic pulse thrombolysis procedure in the treatment of acute submassive pulmonary embolism	No		228			31/05/2017	03/08/2017	04/10/2016	19/03/2018	19/03/2018		D - Sponsor Delays	Sponsor
														H - Contracting delays	
17/LO/0525		A prospective, double-masked, randomized, multicenter, active- controlled, parallel-group, 6-month study assessing the safety and ocular hypotensive efficacy of PG324 Ophthalmic Solution compared to GANFORT® (bimatoprost 0.03%/timolol 0.5%) Ophthalmic Solution in subjects with elevated intraocular pressure (MERCURY 3).	No		37			05/12/2016	25/10/2017	30/06/2017	30/11/2017	01/12/2017	16/01/2018	J - Other	Sponsor
17/LO/0848		A Phase 3, Randomized, Double-Blind, Placebo-Controlled StudyEvaluating the Safety and Efficacy of Selonsertib in Subjects with Nonalcoholic Steatohepatitis (NASH) and Bridging (F3) Fibrosis	No		42				11/10/2017	02/08/2017		22/11/2017	24/11/2017	D - Sponsor Delays	Sponsor
														F - No patients seen	
											ļ			H - Contracting delays	
17/LO/0849		A Phase 3, Randomized, Double-Blind, Placebo-Controlled Study Evaluating the Safety and Efficacy of Selonsertib in Subjects with Compensated Cirrhosis due to Nonalcoholic Steatohepatitis (NASH)	Yes	04/12/2017	43	11	54		11/10/2017	02/08/2017		23/11/2017	24/11/2017		
17/EM/0154		A Multi-centre, Double-blind, Randomised, Placebo-controlled, Parallel-arm Phase IIa Trial to Evaluate the Efficacy, Safety and Tolerability of 28-Day Oral Treatment with PXT002331 (foliglurax) in Reducing Motor Complications of Levodopa Therapy in Subjects with Parkinson's Disease Experiencing End-of-dose Wearing Off and Levodopa- Induced Dyskinesia (AMBLED)	Yes	02/02/2018	4	115	119	08/06/2017	06/10/2017	27/06/2017	08/06/2017	10/10/2017	17/01/2018	D - Sponsor Delays	Sponsor
17/LO/0719	226571		Yes	16/10/2017	31	7	38	20/06/2017	08/09/2017	20/06/2017	04/10/2017	09/10/2017	16/10/2017		

17/EM/0236	221138	Dose finding phase IIb study of Bavisant to evaluate its safety and	No		0			19/06/2017	/ 18/12/2017	08/08/2017	19/06/2017	/ 18/12/2017		H - Contracting delays	Sponsor
		effiCacy in treAtment of exceSsive daytime sleePiness (EDS) in PARkinson's Disease (PD).													
18/LO/0091	140718	Side-Stream Dark Field video analysis of cerebral microcirculation	No		32			12/12/2017	12/02/2018	3 12/02/2018	2	16/03/2018			
10/10/0001	140710	in patients with traumatic brain injury or a brain tumour: the			52			12/12/201/	12/02/2010			10/03/2010			
		effects of blood pressure and intracranial pressure													
17/YH/0296	222070		No					04/09/2017	16/10/2017	7 27/09/2017	7			D - Sponsor Delays	Sponsor
1//10/0290	222979	through wearable sensors and digital alerting.	NO					04/08/2017	10/10/2017	27/09/2017	ĺ			D - Sponsor Delays	Sponsor
17/10/1200	220740		Yes	13/10/2017	35	20		1 12/07/2017	02/08/2017	7 05 /00 /201-	7 07/00/201	07/00/2017	07/00/20	7 J - Other	Noithor
17/LO/1390	229748	-	res	13/10/2017	35	36	7.	1 12/0//201/	03/08/2017	05/09/2017	07/09/2017	07/09/2017	07/09/20	17 J - Other	Neither
		they attend emergency or urgent care with a child under 5 years													
		old in non-urgent situation, in order to reduce future repeat non-													
47/10/4004	220624	urgent attendances	NI					02/07/2017	1 24/00/201-	2 24/00/204	7			D. C	
17/LO/1091	229631	A Phase 3 Clinical Study to Evaluate the Efficacy and Safety of the	NO					03/07/2017	24/08/2017	24/08/2017	′		Sponsor declined site c	on B - Suspended by sponsor	Sponsor
		Combination Regimen of MK-3682B													
		(Grazoprevir/Ruzasvir/Uprifosbuvir) in Participants with Chronic													
		Hepatitis C Virus Genotype 3 Infection													
17/NW/0352	226135		No		159			03/07/2017	05/10/2017	10/08/2017	7	13/03/2018	13/03/20	.8 D - Sponsor Delays	Sponsor
		Trauma													
														H - Contracting delays	
17/SC/0164		A multi-centre, randomised, controlled trial evaluating the effects	Yes	06/09/2017	84	6	90	08/06/2017	08/06/2017	26/05/2017	23/08/2017	31/08/2017	31/08/20	7 E - Staff availability issues	NHS Provi
		of early high-dose cryoprecipitate in adult patients with major													
		trauma haemorrhage requiring major haemorrhage protocol													
		(MHP) activation													
17/EM/0301		Sedation and Weaning in Children	Yes	05/02/2018	16	46	62	2 13/07/2017				21/12/2017			
17/EE/0387			No							8 08/03/2018					
17/LO/1568	232931	Pharmacokinetics of Intramuscular Adrenaline in Food-Allergic	Yes	16/12/2017	26	38	64	4 30/08/2017	13/10/2017	30/10/2017	08/11/2017	08/11/2017		A - Permissions	Neither
		Teenagers - does dose matter? The PIMAT study												delayed/denied	
17/LO/1522	216857	A randomized, double-blind, placebo-controlled, two-cohort	No					31/07/2017	26/03/2018	3 28/11/2017	31/07/2017	'			
		parallel group study to evaluate the efficacy of CAD106 and													
		CNP520 in participants at risk for the onset of clinical symptoms of													
		Alzheimer's disease													
17/LO/1245	208149	A randomised controlled trial to assess the clinical and cost	Yes	08/03/2018	25	31	56	6 11/09/2017	11/01/2018	3 10/11/2017	11/09/2017	05/02/2018			
		effectiveness of topical lactic acid gel for treating second and													
		subsequent episodes of bacterial vaginosis													
17/EE/0026	220207	A Prospective, Randomized, Controlled, Multi-Center Clinical	Yes	05/01/2018	69	85	154	4 15/02/2017	04/08/2017	19/06/2017	21/09/2017	12/10/2017	16/10/20	7 F - No patients seen	Sponsor
		Study of theACRYSOF IQ Extended Depth of Focus (EDF) IOL													
														H - Contracting delays	
17/EM/0361	234065	A Multicenter, Randomized, Double-Blind, Placebo-Controlled	No		14			20/09/2017	28/02/2018	3 11/01/2018	3 20/09/2017	14/03/2018			
		Study in Subjects With Relapsing Multiple Sclerosis to Evaluate the													
		Efficacy and Safety of BIIB033 as an Add-On Therapy to Anti-													
		Inflammatory Disease-Modifying Therapies (AFFINITY)													
16/LO/0831	196728	Efficacy, safety and impact on antimicrobial resistance of duration	No		133			20/06/2017	26/10/2017	/ 11/11/2016	6 08/03/2018	08/03/2018		D - Sponsor Delays	Sponsor
, ,		and dose of amoxicillin treatment with Community-Acquired												, ,	1.
		Pneumonia (CAP): a randomised controlled trial													
17/LO/0380	218003		No		0			25/01/2018	25/01/2018	3 18/05/2017	7 25/01/2018	25/01/2018			
		MULTICENTER, RANDOMIZED, DOUBLE-BLIND, PLACEBO-							,,						
		CONTROLLED, PARALLEL-GROUP, EFFICACY AND SAFETY STUDY OF													
		CRENEZUMAB IN PATIENTS WITH PRODROMAL TO MILD													
		ALZHEIMER'S DISEASE													
17/SC/0582	234/04		No					24/10/2017	24/01/2019	3 18/01/2018	2			D - Sponsor Delays	Sponsor
1/30/0302	234404	characterised oral desensitisation immunotherapy in subjects who	1					24/10/201/	24/01/2010	1 10/01/2010	í l			- Sponsor Delays	5001501
		participated in a prior AR101 study													
		participateu ili a prior Artot study						+		+	+			H. Contracting dalays	
		l								1	1		1	H - Contracting delays	

16/LO/0680	186642	Strategy for maintenance of HIV suppression with	No		39		08/11	/2017	29/01/2018	28/07/2016	09/03/2018	09/03/2018	04/04/2018		
		elvitegravir+darunavir/ritonavir in children (PENTA 17). A two-													
		arm, phase 2/3 multicentre, open-label, randomised study													
		evaluating safety and anti-viral effect of current anti-retroviral													
		therapy compared to elvitegravir (EVG) administered with													
		darunavir/ritonavir (DRV/r) in HIV-1 infected, virologically													
		suppressed paediatric participants													
17/LO/1478	226023	A phase IIIb, open-label, multicentre, international randomised	Yes	08/03/2018	64	14	78 22/09	/2017	20/12/2017	23/11/2017	22/09/2017	22/02/2018		D - Sponsor Delays	Sponsor
		controlled trial of simplified treatment monitoring for 8 weeks													
		glecaprevir (300mg)/pibrentasvir (120mg) in chronic HCV													
		treatment naïve patients without cirrhosis													
														H - Contracting delays	
17/EE/0474	229785	The open-label, randomised, multi-centre, parallel group, two-	No		23		14/07	/2017	10/01/2018	05/01/2018	02/02/2018	02/02/2018	20/02/2018	G - No patients consented	Neither
		arm study to assess the safety, overall tolerability, and antiviral													
		activity of Brincidofovir versus standard of care for treatment of													
		adenovirus infections in high-risk paediatric allogeneic													
		haematopoietic cell transplant recipients													
18/LO/0314	237195	Smart Catheter: A novel biosensor for early diagnosis of catheter	No		14				28/03/2018	06/04/2018		11/04/2018			
		associated urinary tract infection													
17/SC0639	236312	A Phase 3 Open-Label, single-Arm Study to Evaluate The Efficacy	No		80		09/01	/2018	15/01/2018	26/02/2018	22/03/2018	05/04/2018		H - Contracting delays	Sponsor
		and Safety of BMN 270, an Adeno-Associated Virus Vector-													
		Mediated Gene Transfer of Human Factor VIII at a dose of 4E13													
		vg/kg in Haemophila A Patients with Residual FVIII Levels 1 IU/dL													
		Receiving Prophylactic FVIII Infusions													
17/EM/0412	234907	An adaptive seamless randomized, double-blind,	No				15/01	/2018	13/02/2018	04/12/2017	,			D - Sponsor Delays	Sponsor
		placebocontrolled, dose ranging study to investigate the efficacy													
		and safety of LNP023 in primary IgA nephropathy patients													
18/LO/0170	236855	A Phase III, Randomized, Open-label Study to Evaluate	No				29/01	/2018	26/03/2018						
		Pembrolizumab as Neoadjuvant Therapy and in combination with													
		Standard of Care as Adjuvant Therapy for Stage III-IVB Resectable													
		Locoregionally Advanced Head and Neck Squamous Cell													
		Carcinoma (HNSCC)													
18/NW/0031	230387	A multi-centre, randomised, parallel group, open-label, phase II,	No				12/02	/2018	12/02/2018	08/02/2018					
		single-stage selection trial of liposomal irinotecan (nal-IRI) and 5-													
		fluorouracil (5-FU)/folinic acid or docetaxel as second-line therapy													
		in patients with progressive poorly differentiated extra-pulmonary	r												
		neuroendocrine carcinoma (NEC))													
18/SC/0057		A prospective cross-sectional study analysing the utility of novel	No						19/03/2018						
		tablet-based audiometry and an interactive web-based hearing													
		app as screening tools for drug-induced ototoxicity in adults with													
		cystic fibrosis.													
18/LO/0393		A Phase I, Open-Label, Ascending Dose Study to Assess the Safety	No						19/03/2018						
		and Tolerability of AAV2/6 Factor IX Gene Therapy via Zinc Finger													
		Nuclease (ZFN) mediated targeted integration of SB-FIX in Subjects	;												
		with Severe Haemophilia B	ļ								ļ				ļ
16/LO/2150		Randomised Phase II Trial of olaparib, chemotherapy or olaparib	No				20/03	/2018	20/03/2018	26/01/2017					
		and cediranib in patients with BRCA mutated platinum-resistant													
		ovarian cancer													