REC Ref.	IRAS No.	Name of Trial	Target Number	Minimum	Maximum	Target Date	Date	Total	Date That	Total	Reason For Closure Of
NEC NEI.	INAS NO.		Of Patients		Number Of	-	Agreed to	Number Of		Number Of	
			Agreed?	Patients		Patients	recruit	Patients	Closed To	Study	TTa
			Agreeu:	Agreed		Agreed?	target			Participants	
				(Enter	(Enter	Agreeu:	number of	At The	+	Recruited	
				Same In	Same In		patients	Agreed	, c	Neclated	
					Both If Only		patients	Target Date			
				One	One			Target Date			
				Number)	Number)						
16/WS/0005	193858	A PHASE 3, LONG-TERM ACTIVE TREATMENT EXTENSION STUDY OF	Number Agreed		2	Date	01/11/2021	0	21/12/2017	0	Withdrawn By Sponsor
-, -,		MONGERSEN (GED-0301) IN				Agreed	- , , -	-	, , -	-	, , , , , , , , , , , , , , , , , , , ,
		SUBJECTS WITH CROHN?S DISEASE									
15/NE/0389		Phase 1 study to assess Single Doses of ABY-035 in Healthy Subjects and	Number Agreed	3	3	Date	01/02/2018	3	01/02/2018	3	Recruitment Finished
13/102/0305		Psoriasis				Agreed					
		Patients									
16/ES/0001	193859	A Phase 3, randomized, double-blind, placebo-controlled, multicenter study to	Number Agreed	2	2	Date	31/07/2017	1	31/07/2017	1	Recruitment Finished
10, 20, 0001		investigate the efficacy and safety of Mongersen (GED-0301) for the treatment				Agreed					
		of subjects with active Crohn?s disease									
16/LO/0351	190075	A double blind, randomized placebo controlled crossover multiple dose study of	Number Agreed	7	7	Date	16/11/2017	7	07/02/2018	9	Recruitment Finished
		LJN452 to assess safety, tolerability and efficacy in patients with primary bile				Agreed					
		acid diarrhea (pBAD)									
14/NW/1427	166630	A Phase 3, Randomized Study Investigating the Safety of CVT-301 (Levodopa	Number Agreed	4	4	Date	13/12/2017	6	24/07/2017	6	Recruitment Finished
		Inhalation Powder) in Parkinson?s Disease Patients With Motor Response				Agreed					
		Fluctuations (OFF Phenomena) Compared to an Observational Cohort Control									
17/LO/0783	225826	A Multicenter, Parallel-Group, Randomized, Cross-Over Trial to Compare the	Number Agreed	4	4	Date	14/05/2018	0	21/03/2018	0	Withdrawn By Sponsor
		Efficacy of LUMINITY? and SonoVue? in the Evaluation of Left Ventricular				Agreed					
		Endocardial Border Definition									
17/EM/0005	215706	A Double-blind, Randomized, Placebo-controlled, Multicenter Study to Assess	Number Agreed	15	15	Date	05/01/2020	0	29/01/2018	0	Withdrawn By Sponsor
		the Efficacy and Safety of Omecamtiv Mecarbil on Mortality and Morbidity in				Agreed					
		Subjects with Chronic Heart Failure with Reduced Ejection Fraction									
16/SC/0422	209437	A PHASE 2A STUDY OF TRC105 (WITH OPTION TO ADD STANDARD DOSE	Number Agreed	2	2		01/05/2018	0	31/07/2017	0	Withdrawn By Sponsor
		BEVACIZUMAB) IN PATIENTS WITH REFRACTORY GESTATIONAL TROPHOBLASTIC				Agreed					
		NEOPLASIA (GTN)							10/01/0010		
16/NE/0084	196157	A Randomized, Open-label Phase 2 Study of Nanoliposomal Irinotecan (nal-IRI)-	Number Agreed	2	2	Not			18/04/2018	0	Withdrawn By Sponsor
		containing Regimens versus Gemcitabine plus nab-Paclitaxel in Patients with				Available /					
16/YH/0004	100267	Previously Untreated, Metastatic Pancreatic Adenocarcinoma	Denge Agreed	1	2	Not Agreed	20/01/2021	0	07/11/2017	0	Doomuitmoont Finishood
16/YH/0004	190267	A Phase II, Single Arm, Open-Label, Multicenter Study to Evaluate the Efficacy	Range Agreed	1	2	Date	29/01/2021	0	07/11/2017	0	Recruitment Finished
		and Safety of MOR00208 Combined with Idelalisib in Patients with Relapsed or				Agreed					
		Refractory CLL/SLL Previously Treated with Bruton?s Tyrosine Kinase (BTK) Inhibitor									
15/WA/0358	1802///	An exploratory, open-label, multicenter study to evaluate the safety and	Number Agreed	2	2	Date	10/03/2017	2	21/11/2017	3	Recruitment Finished
	105544	efficacy of a two-dose regimen of ATIR101, a T-lymphocyte enriched leukocyte	Agreed	<u></u>		Agreed	10/03/201/	۷	~ 1/ 11/ 201/	5	
		preparation depleted ex vivo of host alloreactive T-cells (using photodynamic				Agreeu					
		treatment), in patients with									
15/NW/0431	1773/16	A Phase 2, Two-arm Multicenter, Open-Label Study to Determine the Efficacy	Number Agreed	2	2	Date	31/12/2017	1	01/12/2017	1	Recruitment Finished
13/100/0431	1//540	and the Safety of Two Different Dose Regimens of a pan-FGFR Tyrosine Kinase		2	<u>_</u>	Agreed	51/12/201/	1	01/12/201/	1	
		Inhibitor JNJ-42756493 in Subjects with Metastatic or Surgically Unresectable				1,5,000					
		Urothelial Cancer with FGFR									
	ļ		ļ	I	1	I			1		

16/EM/0259	165592 A prospective, multicentre, open-label, randomized, active-controlled, 3 parallel	Number Agreed	2	2	Date	22/07/2017	3	22/07/2017	3	Recruitment Finished
	groups, phase 2 study to compare the efficacy and safety of masitinib in combination with FOLFIRI (irinotecan, 5-fluorouracil and folinic acid), versus				Agreed					
	masitinib alone, versu									
15/WM/0327	185434 Clinical evaluation of the Therapeutic IntraVascular Ultrasound (TIVUS?) System	Number Agreed	5	5	Date	30/06/2017	2	02/10/2017	2	Recruitment Finished
14/50/0202	for pulmonary artery denervation in patients with pulmonary hypertension		6	C	Agreed	20/10/2017	9	14/00/2017	9	Doorwitze out Finished
14/SC/0262		Number Agreed	D		Date	30/10/2017	9	14/09/2017	9	Recruitment Finished
	with neovascular Age-related macular degeneration (nAMD), evaluating the				Agreed					
	efficacy and safety of switching From intravitreal Aflibercept to RanIbizumab 0.5mg: the SAFARI study									
16/WM/0271		Number Agreed	10	10	Date	07/07/2021	5	27/10/2017	5	Withdrawn By Sponsor
10, 00101, 02, 1	Neurovascular System?				Agreed				-	
16/EM/0180		Number Agreed	8	8	Date	21/09/2017	3	29/08/2017	3	Withdrawn By Sponsor
	Study to Select the Daily Oral Dose of Estetrol (E4) for the Treatment of	-			Agreed					
	Vasomotor Symptoms in Post-Menopausal Women									
09/H0724/39	Evaluation of Safety and Effectiveness of the FormulaTM PTMTM Balloon-	Not Available /			Not			03/10/2017	0	Recruitment Finished
	Expandable stent for Renal Artery Stenosis	Not Agreed			Available /					
					Not Agreed					
16/EE/0463	214371 An open-label, multicenter, Phase IIIb study to assess the safety and efficacy of	Number Agreed	1	1	Date	31/10/2017	1	07/11/2017	1	Recruitment Finished
	ribociclib (LEE011) in combination with letrozole for the treatment of men and				Agreed					
	pre/postmenopausal women with hormone receptor-positive (HR+) HER2-									
	negative (HER2-) advanced br									
16/WM/0433	215494 A research study to describe the ?real world? use of Thrombopoietin-Receptor	Number Agreed	20	20	Date	31/08/2017	39	31/08/2017	39	Recruitment Finished
	Agonists (TRAs) in the management of Immune Thrombocytopenia (ITp) in the				Agreed					
	UK									
17/LO/0100		Number Agreed	3	3	Date	31/05/2018	0	22/09/2017	0	Withdrawn By Sponsor
	Available Therapy for the Treatment of Severe Infections Caused by				Agreed					
	Carbapenem-resistant Gram-negative Pathogens									
16/WM/0250	194366 A prospective, multi-national, multicentre, non-interventional study to evaluate	Number Agreed	10	10	Date	31/03/2018	4	31/03/2018	4	Withdrawn By Sponsor
	the long term safety of Esmya, in particular the endometrial safety, and the				Agreed					
	current prescription and management patterns of Esmya in a long term									
16/10729	treatment setting.	Not Available /			Not			21/12/2017	0	Recruitment Finished
16/NW/0728		Not Available / Not Agreed			Not Available /			31/12/2017	0	Recruitment Finished
	development of good sleep habits in bables under 12 months of age	NOL Agreeu			Not Agreed					
16/EM/0376	211430 The LEADERSHIP 301 Trial: A 12-Week, Randomized, Multi-Center, Double-Blind,	Number Agreed	04/01/1900			15/02/2018	0	15/02/2018	0	Recruitment Finished
10/ 2101/ 03/ 0	Placebo-Controlled, 3-Arm, Parallel-Group, Phase 3 Trial to Evaluate the Efficacy	Number Agreed	04/01/1500	04/01/1500	Agreed	15/02/2018	0	13/02/2018	0	Reci ditilient i misned
	and Safety of 2 Doses of AQX-1125 Targeting the SHIP1 Pathway in Subjects				, greed					
	with Interstitial Cysti									
16/SS/0134	199690 A PHASE 1B MULTICENTER, OPEN-LABEL STUDY TO EVALUATE THE SAFETY AND	Number Agreed	03/01/1900	03/01/1900	Date	28/02/2019	0	02/03/2018	0	Withdrawn By Sponsor
10,00,010 .	TOLERABILITY AND DETERMINE THE MAXIMUM TOLERATED DOSE OF PF-	U			Agreed					, ,
	05230907 IN SUBJECTS									
15/LO/2098	182147 A Phase III Double-blind, Randomised, Parallel-Group Comparison of the Efficacy	Number Agreed	14/01/1900	14/01/1900	Date	01/02/2018	3	11/12/2017	3	Recruitment Finished
	and Safety of FP-1201-lyo (Recombinant Human Interferon Beta-1a) and	-			Agreed					
	Placebo in the Treatment of Patients with Moderate or Severe Acute									
	Respiratory Distress Syndrome									
14/NI/0002	143644 AN OPEN LABEL, REGISTRY STUDY OF THE SAFETY OF ILUVIEN? 190	Number Agreed	12/01/1900	12/01/1900	Date	30/12/2016	12	12/07/2017	12	Recruitment Finished

07/H1102/84		Sunitinib treatment of renal adjuvant cancer (S-TRAC) A randomised double blind phase 3 study of adjuvant vs placebo in subjects at high risk of recurrent	Number Agreed	03/01/1900	03/01/1900	Not Available /		30/08/20)17 (Recruitment Finished
		RCC. Study Number: A6181109				Not Agreed				
12/YH/0179	102439	A Companion Trial to Study GS-US-312-0116: A Phase 3, Randomized, Double-	Number Agreed	02/01/1900	02/01/1900		19/11/2017	2 15/02/20)18 2	Recruitment Finished
12, 11, 01, 5	102455	Blind, Placebo-Controlled Study Evaluating the Efficacy and Safety of GS-1101	interinser Agreed			Agreed	15/11/2017	2 13/02/20	2	
		(CAL-101) in Combination with Rituximab as Therapy for Patients with				, gi ccu				
		Previously Treated Chronic Lymphocyt								
17/NW/0175	222850	A Phase 1b Open-Label, Dose Escalation Study of PRTX-100 in Adult patients	Number Agreed	01/01/1000	01/01/1900	Date	20/12/2017	2 17/05/20	119 2	Recruitment Finished
1//11/01/5	222033	with Persistent/Chronic Immune Thrombocytopenia	Number Agreeu	01/01/1900			20/12/2017	2 17/05/20		
17/LO/0304	222475	A double-blind, randomised, placebo-controlled, parallel-group, phase 2, dose-	Range Agreed	10/01/1000	12/01/1900	Agreed	31/10/2018	0 31/08/20)17 (Withdrawn By Sponso
17/10/0304	222475		Ralige Agreeu	10/01/1900	12/01/1900		51/10/2018	0 51/06/20		withurawin by sponso
		ranging trial to evaluate the efficacy, safety and tolerability of oral Litoxetine				Agreed				
		10mg, 20mg and 40mg twice daily (BID) versus placebo in women with mixed								
47/55/0004	240.405	urinary continence		02/04/4000	02/04/4000	Data	04/05/2020	17 01 /05 /06	10 1-	
17/EE/0081	219405	An open label, single arm pilot study of OncoSil?, administered to study	Number Agreed	02/01/1900	02/01/1900		01/06/2020	17 01/06/20	18 17	Recruitment Finished
		participants with unresectable locally advanced pancreatic adenocarcinoma,				Agreed				
		given in combination with FOLFIRINOX or gemcitabine+nab-paclitaxel								
		chemotherapies.								
16/LO/2148	219354	A Multi-Centre, Open Label, Phase IIb Study, Evaluating the Safety, Tolerability	Number Agreed	04/01/1900	04/01/1900	Date	01/01/2018	8 01/01/20	18 8	Recruitment Finished
		and Efficacy of Targeted Intraprostatic Administration of PRX302 to Treat Men				Agreed				
		with Histologically Proven, Clinically Significant, Localised, Low- to Intermediate-								
		Risk Prostate								
16/WM/0006	192580	WHIST - Wound Healing In Surgery for Trauma. A Randomised Controlled Trial	Number Agreed	11/02/1900	11/02/1900	Date	30/04/2018	37 30/04/20)18 37	Recruitment Finished
		of standard wound management				Agreed				
		versus negative pressure wound therapy in the treatment of adult patients								
		having surgical incisions for major trauma								
		to the lower limb								
14/LO/0344	140294	A Phase 3, Open-Label, Randomized, Parallel, 2-Arm, Multi-Center	Number Agreed	01/01/1900	01/01/1900	Date	01/07/2017	2 01/07/20)17 2	Recruitment Finished
		Study of Talazoparib (BMN 673) versus Physician?s Choice in				Agreed				
		Germline BRCA Mutation Subjects with Locally Advanced and/or								
		Metastatic Breast Cancer, Who Have Received Prior Chemotherapy Regimen								
16/YH/0022	194494	A phase I, multicenter, open-label, single-sequence drugdrug interaction study	Number Agreed	03/01/1900	03/01/1900	Date	16/08/2016	0 05/07/20)17 (Recruitment Finished
		to assess the effect of INC280 on the pharmacokinetics of midazolam and				Agreed				
		caffeine in patients with cMET-dysregulated advanced solid tumors								
16/EE/0009	189564	A Phase 3,Randomized,Controlled, Multi-Center,Open-Label,Study To Compare	Number Agreed	03/01/1900	03/01/1900	Not		11/08/20)17 1	Withdrawn By Sponso
		Tivozanib Hydrochloride To Sorafenib In Subjects With Refractory Advanced				Available /				, , ,
		Renal Cell Carcinoma				Not Agreed				
15/LO/1232	182226		Number Agreed	03/01/1900	03/01/1900	-	30/11/2017	7 22/02/20)18 7	Recruitment Finished
10, 10, 1202	102220	Combination as an Adjunctive Therapy to a Prostaglandin Analogue		00,01,1500		Agreed	00,11,201,	,, 02, 20	,10	
16/YH/0081	186580	A Multicentre, Randomised, Double-blind, Placebo-controlled, Phase 2 Study	Number Agreed	01/01/1900	01/01/1900	-	24/04/2018	0 24/04/20)18 (Recruitment Finished
10/11/0001	100500	Evaluating the Efficacy and Safety of Anifrolumab in Adult Subjects with Active	intering ceu			Agreed	24/04/2010	0 24/04/20		
		Proliferative Lupus Nephritis.				Agreeu				
16/NE/0023	100025	A Phase III, Multi-centre, Randomized, Double-Blind, Placebo-Controlled,	Number Agreed	08/01/1000	08/01/1000	Date	01/08/2018	8 01/08/20	17 9	Recruitment Finished
10/ NE/ 0025	120022	Parallel-Group, Efficacy and Safety Study with Open-Label Extension of	Inditibel Agreed	00/01/1900			01/00/2010	01/00/20	···/ c	
						Agreed				
16/10/0124	102055	Crenezumab in Patients with Prodromal-to-Mild Alzheimer?s Disease	Number	10/01/1000	10/01/1000	Data	17/12/2010	10 17/10/00	17 40	
16/LO/0124	182922	A study of the ReCor Paradise System in Clinical Hypertension	Number Agreed	10/01/1900	10/01/1900		17/12/2019	10 17/12/20	10	Recruitment Finished
	400000			40/04/1100-	40/04/1000	Agreed			47	
15/LO/1538	188226	A study to evaluate the effectiveness and safety of Exablate transcranial	Number Agreed	10/01/1900	10/01/1900		13/02/2019	0 16/10/20	01/ 0	Recruitment Finished
		MRgFUS thalamotomy treatment of medication refractory essential tremor subjects				Agreed				

15/LO/1641	180563	Global Clinical Study of Renal Denervation with the Symplicity Spyral? multielectrode renal denervation system in Patients with Uncontrolled Hypertension in the	Number Agreed	05/01/1900	05/01/1900	Date Agreed	01/11/2019	6 04/06/2018	6 Recruitment Finished
15/LO/1163	181430	Absence of Antihypertensive Medications A Phase 3b, Randomized, Double-Blind, Switch Study to Evaluate F/TAF in HIV-1 Infected Subjects who are Virologically Suppressed on Regimens containing ABC/3TC	Number Agreed	30/01/1900	30/01/1900	Date Agreed	31/12/2017	5 30/03/2018	5 Recruitment Finished
15/LO/1640	180561	Global Clinical Study of Renal Denervation with the Symplicity Spyral? multielectrode renal denervation system in patients with uncontrolled hypertension on standard medical therapy	Number Agreed	05/01/1900	10/01/1900	Date Agreed	31/12/2019	1 03/05/2018	1 Withdrawn By Host
15/LO/0141	169126	The Efficacy and Safety of Bimatoprost SR in Patients With Openangle Glaucoma or Ocular Hypertension	Number Agreed	01/01/1900	01/01/1900	Date Agreed	30/11/2017	1 30/11/2017	1 Recruitment Finished
15/LO/1487	172859	A Multicenter, Randomized, Double-Blind Study of Erlotinib in Combination with Ramucirumab or Placebo in Previously Untreated Patients with EGFR Mutation- Positive Metastatic Non-Small Cell Lung Cancer	Number Agreed	03/01/1900	03/01/1900	-	01/08/2018	3 01/06/2018	3 Recruitment Finished
15/EM/0021	166923	A randomised, double blind, placebo controlled multicenter trial, examining the effect of Natrox? on the rates of healing for chronic diabetic foot ulcers	Number Agreed	05/01/1900	05/01/1900	Date Agreed	01/08/2017	7 01/08/2017	7 Recruitment Finished
14/LO/1728	158859	A Phase Ib Open Label Study of the Safety and Tolerability of ME-344 Given in Combination with Topotecan (Hycamtin?) in Patients with Solid Tumors	Number Agreed	02/01/1900	02/01/1900	Date Agreed	31/07/2017	2 31/07/2017	2 Recruitment Finished
14/LO/2143	164828	Measurement of low-energy stimulation in patients with atrial fibrillation	Number Agreed	10/01/1900	10/01/1900	Date Agreed	31/08/2017	3 31/08/2017	3 Recruitment Finished
11/EE/0256		Multi-centre, open label, prospective, consecutive series registry database of BioPoly RS Partial Resurfacing Knee implant	Number Agreed	10/01/1900	10/01/1900	Date Agreed	31/12/2016	0 01/06/2018	0 Recruitment Finished
12/NW/0137	98327	A randomized, double-blind, multicenter, Phase III study of everolimus (RAD001) plus best supportive care versus placebo plus best supportive care in the treatment of patients with advanced NET of GI or lung origin	Number Agreed	05/01/1900	05/01/1900	Date Agreed	03/11/2017	5 03/11/2017	5 Recruitment Finished
16/EE/0133	202599	Safety and proof of principle study of ATX-GD-59 in male and female subjects with Graves? disease not currently treated with anti-thyroid therapy: An Open label study, with an upward titration over five dose levels administered by Intradermal injection	Range Agreed	03/01/1900	05/01/1900	Date Agreed	30/01/2018	1 30/01/2018	1 Recruitment Finished
16/LO/0886	204525	A Multicenter, Double-Blind, Placebo-Controlled, Randomized, Parallel-Group, Dose-Ranging Study to Evaluate the Safety and Efficacy of Intravenous Natalizumab (BG00002) in Acute Ischemic Stroke	Number Agreed	03/01/1900	03/01/1900	Date Agreed	31/08/2017	7 31/08/2017	7 Recruitment Finished
14/WS/1146	166537	Multicenter, randomized, double-blind, double-dummy, active-comparator, event-driven, superiority phase III study of secondary prevention of stroke and prevention of systemic embolism in patients with a recent Embolic Stroke of Undetermined Source (ESUS),	Number Agreed	20/01/1900	20/01/1900	Date Agreed	31/12/2016	20 15/01/2018	27 Recruitment Finished
17/LO/0082	217324	A Multi-center, Single Arm Post-market Clinical Study to Confirm Safety and Performance of PuraStat Absorbable Haemostatic Material for the Management of Bleeding in Vascular Surgery	Number Agreed	03/01/1900		Date Agreed	31/12/2017	4 31/12/2017	4 Recruitment Finished
16/LO/0118	191634	Valiant Evo International Clinical Trial	Number Agreed	04/01/1900	04/01/1900	Date Agreed	30/07/2018	5 01/06/2018	5 Recruitment Finished
17/WS/0030		A Phase Ib and II Open-Label, Multi-Center Study of MEDI4736 Evaluated in Different Combinations in Patients with Metastatic Pancreatic Ductal Adenocarcinoma	Number Agreed	02/01/1900	02/01/1900	-	30/11/2017	4 03/03/2018	4 Recruitment Finished