Researc h Ethics Committ ee Referen ce Number	Integrated Research Applicatio n System Number	Name of Trial	First Participa nt Recruite d?	Date of First Participant Recruited	Date Site Invited	Date Site Selected	HRA Approval Date	Date Site Confirmed By Sponsor	Date Site Confirmed	Non- Confirmation Status	Date Site Ready To Start	Reasons for Delay	Reasons for delay correspond to:
18/YH/0 417	247000	Phase 1 Multiple-Ascending-Dose Study to Assess the Safety, Tolerability, and Pharmacokinetics of BIIB078 Administered Intrathecally to Adults with C9ORF72- Associated Amyotrophic Lateral Sclerosis	No		29/11/2018	29/11/2018				Site declined to participate			Please Select
18/EM/ 0229	250101	A RANDOMIZED, MULTICENTER, DOUBLE-BLIND, PARALLEL-GROUP, ACTIVE-CONTROL STUDY OF THE EFFICACY AND SAFETY OF SPARSENTAN FOR THE TREATMENT OF IMMUNOGLOBULIN A NEPHROPATHY	Yes	16/04/2019	16/08/2018	08/10/2018	04/08/2018	12/11/2018	20/12/2018	Please Select	16/01/2019	D - Sponsor Delays	Sponsor
18/WA/ 0154	233884	WHITE 8 COPAL: A Randomised Controlled Trial of low dose single antibiotic loaded cement versus high dose dual antibiotic loaded cement in patients receiving a hip hemiarthroplasty after fracture	Yes	26/04/2019	10/09/2018	21/11/2018	10/05/2018	28/02/2019	09/04/2019	Please Select	09/04/2019		Please Select
17/LO/0 871	215297	Methylphenidate versus placebo for fatigue in advanced cancer (MePFAC)	Yes	30/01/2019	28/08/2018	29/11/2018	23/05/2018	03/01/2019	09/01/2019	Please Select	10/01/2019		Please Select
18/LO/2 067	253346	A Phase 2 Study of INCMGA00012 in Participants With Squamous Carcinoma of the Anal Canal Who Have Progressed Following Platinum-Based Chemotherapy	Yes	19/04/2019	17/12/2018	17/12/2018	07/01/2019	11/03/2019	20/03/2019	Please Select	09/04/2019		Please Select
18/LO/0 555	237992	Preventing Ovarian Cancer through early Excision of Tubes and late Ovarian Removal	Yes	19/06/2019	28/01/2019	21/03/2019	19/04/2018	20/03/2019	20/05/2019	Please Select	20/05/2019		Please Select
17/YH/0 120	208838	A pragmatic multi-centre randomised controlled non- inferiority, cost effectiveness trial comparing injections of collagenase into the cord to surgical correction in the treatment of moderate Dupuytren's Contracture in adult patients	Yes	17/05/2019	21/12/2018	21/12/2018	25/05/2017	15/04/2019	17/04/2019	Please Select	17/04/2019		Please Select
18/WA/ 0092	230113	USTEKID – Phase II multi-centre, double-blind, randomised trial of Ustekinumab in adolescents with new-onset type 1 diabetes	Yes	30/05/2019	02/01/2019	02/01/2019	21/09/2018	11/02/2019	30/04/2019	Please Select	01/05/2019		Please Select

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18/EM/ 0193	243749	A RANDOMIZED, MULTICENTER, DOUBLE-BLIND, PARALLEL, ACTIVE-CONTROL STUDY OF THE EFFECTS OF SPARSENTAN, A DUAL ENDOTHELIN RECEPTOR AND ANGIOTENSIN RECEPTOR BLOCKER, ON RENAL OUTCOMES IN PATIENTS WITH PRIMARY FOCAL SEGMENTAL GLOMERULOSCLEROSIS	No		15/01/2019	19/02/2019	18/09/2018	26/03/2019	24/04/2019	Please Select	08/05/2019	F - No patients seen	Neither
18/LO/0 324	220073	The VIDEO Trial: A Feasibility RCT of Vitrectomy Plus Standard Care Intravitreal Ranibizumab Injections versus Standard Care Intravitreal Ranibizumab injections Alone In Patients With Centre Involving Diabetic Macular Edema	Yes	01/05/2019	26/03/2019	26/03/2019	03/04/2019	26/03/2019	26/03/2019	Please Select	26/03/2019	D - Sponsor Delays	Sponsor
18/LO/1 453	246119	BARBICAN: A randomised, open-label Phase II study to determine the contribution of ipatasertib to neoadjuvant chemotherapy plus atezolizumab in women with triple-negative breast cancer	No		27/03/2019	27/03/2019	15/11/2018			Please Select		H - Contracting delays	Sponsor
19/YH/0 015	255446	Beta Blockers or placebo for primary prophylaxis of oesophageal varices (BOPPP Trial). A blinded, multi- centre, clinical effectiveness and cost-effectiveness randomised controlled trial	Yes	30/08/2019	27/03/2019	27/03/2019	24/04/2019	23/05/2019	17/06/2019	Please Select	21/06/2019	D - Sponsor Delays	Sponsor
17/EE/0 368	213669	A randomised trial to assess whether the addition of a beta blocker infusion (landiolol) to standard treatment in patients with septic shock, requiring prolonged (>24 hours) support with high-dose vasopressor agents, improves organ failure (the STRESS-L trial)	No		10/01/2019	03/05/2019	10/11/2017			Please Select			Please Select
19/YH/0 151	257273	B7981015 - PF-06651600 IN ADULT AND ADOLESCENT ALOPECIA AREATA (AA)	No		09/11/2018	09/05/2019	24/07/2019	11/09/2019	16/09/2019	Please Select		D - Sponsor Delays	Sponsor
19/SW/ 0093	260867	A Phase 2, Randomized, Double-blind, Placebo- controlled Evaluation of the Safety and Efficacy of BMS-986165 with Background Treatment in Subjects with Lupus Nephritis	No		25/04/2019	24/05/2019	17/06/2019	25/04/2019	04/09/2019	Please Select		D - Sponsor Delays	Sponsor
19/SC/0 021	249552	OPtimal TIMing of Anticoagulation after acute ischaemic Stroke: a randomised controlled trial (OPTIMAS Trial)	No		13/06/2019	13/06/2019	04/04/2019			Please Select			Please Select
19/LO/0 905	257865	A Phase IV, Randomised, Open-Label Pilot Study to Evaluate Switching from Protease-Inhibitor based regimen to Bictegravir/Emtricitabine/Tenofovir Alafenamide Single Tablet Regimen in Integrase Inhibitor-naïve, virologically suppressed HIV-1 infected adults harbouring drug resistance mutations.	No		17/05/2019	28/06/2019	17/07/2019	05/09/2019	16/09/2019	Please Select	16/09/2019		Please Select

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19/LO/0 496	259483	A Phase 3b Multicenter, Randomized, Double-Blind, Double-Dummy, Active Controlled Study Comparing the Safety and Efficacy of Upadacitinib to Dupilumab in Adult Subjects with Moderate to Severe Atopic Dermatitis	No		19/12/2018	15/01/2019	10/06/2019	24/07/2019	13/09/2019	Please Select	13/09/2019		Please Select
18/LO/1 187	240011	A Multicenter, Randomized, Double-Blind, Placebo- Controlled Phase 3 Study in Ovarian Cancer Patients Evaluating Rucaparib and Nivolumab as Maintenance Treatment Following Response to Front-Line Platinum- Based Chemotherapy	Yes	08/07/2019	16/10/2018	21/01/2019	11/09/2018	21/03/2019	21/03/2019	Please Select	11/04/2019	G - No patients consented	Neither
18/NW/ 0682	249138	Clinical and Cost-effectiveness of Posterior Cervical Foraminotomy versus Anterior Cervical Discectomy in the Treatment of Cervical Brachialgia: A Multicentre, Phase III, Randomised Controlled Trial	No		19/02/2019	19/02/2019	19/11/2018			Please Select			Please Select
19/LO/0 040	253233	A PHASE 4, MULTI-CENTER, RANDOMIZED, DOUBLE- BLIND, PLACEBO-CONTROLLED STUDY OF THE IMPACT OF APREMILAST (CC-10004) ON QUALITY OF LIFE, EFFICACY, AND SAFETY IN SUBJECTS WITH MANIFESTATIONS OF PLAQUE PSORIASIS AND IMPAIRED QUALITY OF LIFE	No		04/03/2019	04/03/2019	15/08/2018	25/04/2019	15/05/2019	Please Select	15/05/2019	D - Sponsor Delays F - No patients seen	Sponsor
19/EM/ 0034	255895	A multicentre, randomised, double-blind, placebo- controlled, parallel-group study to evaluate the efficacy and safety of padsevonil as adjunctive treatment of focal-onset seizures in adult subjects with drug-resistant epilepsy	No		15/05/2019	16/09/2019	16/06/2019	01/10/2019	01/10/2019	Please Select			Please Select
19/LO/0 166	258220	The VITALE Study Evaluating Safety and Effectiveness/Performance of the Microport CardioFlow VitaFlow II - Transcatheter Aortic Valve System. VitaFlowTM II Transcatheter Aortic Valve System Study	No		28/03/2019	15/07/2019	11/04/2019	25/09/2019		Please Select			Please Select
19/LO/0 738	258589	A randomised, two-arm (1:1 ratio), double blind, placebo controlled phase III trial to assess the efficacy, safety, cost and cost-effectiveness of rituximab in treating de novo or relapsing NS in patients with MCD/FSGS (TURING)	No		17/07/2019	17/07/2019	14/06/2019			Please Select			Please Select

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18/WM /0394	248493	PErioperAtive CHildhood obesitY (PEACHY): A prospective observational cohort study investigating the proportion of overweight and obese children presenting for a procedure under general anaesthesia in the UK and the incidence of preoperative adverse outcomes in this patient group	Yes	10/09/2019	23/08/2019	23/08/2019	13/02/2019	10/09/2019	10/09/2019	Please Select	10/09/2019		Ple
17/SC/0 434	209815	Biomarker-guided duration of antibiotic treatment for sepsis	No		25/04/2019	26/04/2019	23/10/2017			Please Select			Please Select
18/LO/1 674	244500	Randomised controlled trial of an app-based digital intervention to support breast cancer survivors prescribed hormone therapy (e-path study)	Yes	09/10/2019	17/06/2019	31/07/2019	17/11/2018	30/09/2019	30/09/2019	Please Select	30/09/2019		Please Select
18/SC/0 624	244229	Short or Long Antibiotic Regimes in Orthopaedics (SOLARIO): A Randomised Open Label Multi-Centre Clinical Trial	No		22/08/2019	22/08/2019	21/12/2018			Please Select			Please Select
17/YH/0 311	229294	A Modular, Multipart, Multiarm, Open-label, Phase I/lla Study to Evaluate the Safety and Tolerability of CT7001 Alone and in Combination with Anti-cancer Treatments in Patients with Advanced Malignancies	No		19/06/2019	10/09/2019	30/10/2017			Please Select			Please Select
19/LO/1 569	270111	A phase 3b, open label treatment extension study of upadacitinib for the treatment of adult subjects with moderate to severe atopic dermatitis who successfully completed treatment in the M16-046 study	No		24/09/2019	24/09/2019				Please Select			Please Select
19/LO/0 802	264414	Acceptability and tolerance study of high energy peptide supplement	No		29/05/2019	29/05/2019	21/05/2019			Please Select			Please Select
18/WA/ 0199	108978	A randomised, placebo controlled trial of azithromycin for the prevention of chronic lung disease of prematurity in preterm infants	No		08/05/2019	18/07/2019	29/06/2018			Please Select			Please Select
19/SC/0 507	265282	RESPOND EDGE: Repositionable Lotus Edge™ Valve System – Post Market Evaluation of Real World Clinical Outcomes	No		30/09/2019	30/09/2019				Please Select			Please Select
19/EM/ 0220	265213	A Phase III, randomized, multicenter, open-label, non- inferiority study evaluating the efficacy, safety and tolerability of switching to dolutegravir/lamivudine fixed dose combination in HIV-1 infected adults who are virologically suppressed	No		22/08/2019	24/09/2019	11/09/2019			Please Select			Please Select
19/WM /0219	261627	High Flow humidified oxygen as an early intervention in children with Acute Severe Asthma. A feasibility study (HiFlo ASA)	No		02/09/2019	02/09/2019	08/08/2019			Please Select			Please Select
19/LO/1 217	264929	Watchman FLX Left Atrial Appendage Closure Device Post Approval Study	No		05/09/2019	05/09/2019	04/09/2019			Please Select			Please Select