Brighton and Sussex University Hospitals NHS Trust Clinical Trials Initiation Performance Q1 2020 to 2021

Research Ethics Committ e Referenc e Number	Integrate d Research Applicati on System Number	Name of Trial	First Participa nt Recruite d?	Date of First Participant Recruited	Duration between Date Site Selected and Date Site Confirme d	Duration between Date Site Confirme d and First Participa nt Recruite d	Duration between Date Site Selected and First Participa nt Recruite d	Date Site Invited	Date Site Selected	HRA Approval Date	Date Site Confirmed By Sponsor	Date Site Confirmed	Non- Confirma tion Status	Date Site Ready To Start	Reasons for Delay	Comments	Reasons for delay correspo nd to:
19/EM/00 34	255895	A multicentre, randomised, double-blind, placebo-controlled, parallel-group study to evaluate the efficacy and safety of padsevorial as adjunctive treatment of focal- onset seizures in adult subjects with drug- resistant epilepsy	Yes	03/12/2019	15	63	78	15/05/2019	16/09/2019	16/06/2019	01/10/2019	01/10/2019	Please Select	23/10/2019		Contract signed electronically in error by host site too early	NHS Provider
19/LO/01 66	258220	The VITALE Study Evaluating Safety and Effectiveness/Performance of the Microport CardioFlow VitaFlow II - Transcatheter Aortic Valve System. VitaFlowTh II Transcatheter Aortic Valve System Study	Yes	21/11/2019	128	1	129	28/03/2019	15/07/2019	11/04/2019	25/09/2019	20/11/2019	Please Select	20/11/2019		Initiation target met	Please Select
19/LO/07 38	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)	Yes	04/03/2020	223	8	231	17/07/2019	17/07/2019	14/06/2019	06/02/2020	25/02/2020	Please Select	25/02/2020		Initiation target met	Please Select
18/WM/0 394	248493	PErioperAtive CHildhood obesitY (PEACHY): A prospective observational cohort study investigating the proportion of overveight and obese children presenting for a procedure under general anaesthesis in the UK and the incidence of preoperative adverse outcomes in this patient group	Yes	10/09/2019	18	0	18	23/08/2019	23/08/2019	13/02/2019	10/09/2019	10/09/2019	Please Select	10/09/2019		Initiation target met	Please Select
18/LO/16 74	244500	Randomised controlled trial of an app-based digital intervention to support breast cancer survivors prescribed hormone therapy (epath study)	Yes	14/10/2019	61	14	75	17/06/2019	31/07/2019	17/11/2018	30/09/2019	30/09/2019	Please Select	30/09/2019		Initiation target met	Please Select
18/SC/06 24	244229	Short or Long Antibiotic Regimes in Orthopaedics (SOLARIO): A Randomised Open Label Multi-Centre Clinical Trial	Yes	21/11/2019	24	13	37	22/08/2019	15/10/2019	21/12/2018	08/11/2019	08/11/2019	Please Select	08/11/2019		Initiation target met	Please Select
17/YH/03 11	229294	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	No		149			19/06/2019	10/09/2019	30/10/2017	12/11/2019	06/02/2020	Please Select	06/02/2020	F - No patients seen	Study temporarily closed due to COVID 19 and then no eligible pts seen	Neither
18/WA/0 199	108978	A randomised, placebo controlled trial of azithromycin for the prevention of chronic lung disease of prematurity in preterm infants	No		230			08/05/2019	18/07/2019	29/06/2018	26/02/2020	04/03/2020	Please Select	04/03/2020	J - Other	Study recruitment was suspended due to COVID 19	Neither

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19/SC/05 07	265282	RESPOND EDGE: Repositionable Lotus Edge™ Valve System – Post Market Evaluation of Real World Clinical Outcomes	Yes	06/01/2020	85	13	98	30/09/2019	30/09/2019	12/11/2019	05/12/2019	24/12/2019	Please Select	24/12/2019		Initiation target met	Please Select
19/EM/02 20	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	Yes	02/12/2019	74	31	105	19/08/2019	19/08/2019	11/09/2019	11/10/2019	01/11/2019	Please Select	01/11/2019		Initiation target met	Please Select
19/WM/0 219	261627	High Flow humidified oxygen as an early intervention in children with Acute Severe Asthma. A feasibility study (HiFlo ASA)	Yes	23/02/2020	168	6	174	02/09/2019	02/09/2019	08/08/2019	17/02/2020	17/02/2020	Please Select	17/02/2020		Initiation target met	Please Select
19/LO/12 17	264929	Watchman FLX Left Atrial Appendage Closure Device Post Approval Study	Yes	20/11/2019	71	6	77	28/05/2019	04/09/2019	04/09/2019	06/11/2019	14/11/2019	Please Select	14/11/2019		Initiation target met - same day as sponsor green light!	Please Select
17/NW/0 581	214739	Doss Interleukin-1 Receptor Antagonist Improve Outcome following aneurysmal Subarachnoid Haemorrhage (aSAH)? A Phase III trial	Yes	17/10/2019	64	30	94	14/08/2018	15/07/2019	17/06/2019	06/11/2018	17/09/2019	Please Select	17/09/2019	H - Contracti ng delays	10 days between CTA signature / C&C and Green Light. The Sponsor wouldn's repeat of the Community of the Com	Both
19/LO/02 55	251219	LITTLE JOURNEY: A multi-centre randomised controlled trial assessing the effectiveness of the Little Journey app at reducing peri-operative anxiety compared to standard care	No		133			04/09/2019	05/09/2019	08/05/2019	16/01/2020	16/01/2020	Please Select	27/01/2020	J - Other	Recruitment on hold - COVID 19	Neither
19/NW/0 158	259931	SCIENCE Surgery or Cast for Injuries of the EpicoNdyle in Children's Elbows:A multi- centre prospective randomised superiority trial of operative fixation versus non-operative treatment for medial epicondyle fractures of the humerus in children.	Yes	09/12/2019	94	0	94	10/05/2019	06/09/2019	25/03/2019	25/11/2019	09/12/2019	Please Select	09/12/2019		Initiation target met	Please Select
16/LO/13 18	187932	Nucleos(t)ide withdrawal in HBeAg negative hepatitis B virus infection to promote HBsAg clearance. (NUC-B)	No		101			12/09/2018	18/10/2019	21/09/2016	16/01/2020	27/01/2020	Please Select	17/02/2020	J - Other	Recruitment suspended due to COVID 19	Neither
19/SW/0 154	265849	Visual and Optical outcomes after bilateral implantation of Tecnis Eyhance versus Rayner RayOne aspheric in patients undergoing routine cataract surgery	Yes	13/11/2019	11	12	23	18/10/2019	21/10/2019	23/10/2019	01/11/2019	01/11/2019	Please Select	01/11/2019		Initiation target met	Please Select
15/WA/0 395	266296	A randomised Phase II/III trial to study radiotherapy dose escalation in patients with ossophageal cancer treated with definitive chemo-radiation with an embedded Phase II trial for patients with a poor early response using positron emission tomography (PET)	Yes	24/01/2020	188	8	196	11/01/2019	12/07/2019	31/10/2016	16/01/2020	16/01/2020	Please Select	16/01/2020		Initiation target met	Please Select
20/NW/0 035	272039	A Phase 3, Randomized, Double-blind, Multicenter Study of Dostarlimab (TSR-042) plus Carboplatin-Pacifizate Versus Placebo plus Carboplatin-Pacifizate in Pasients with Recurrent or Primary Advanced Endometrial Cancer (RUBY)	No		285			28/11/2019	18/12/2019	20/03/2020	01/07/2020	28/09/2020	Please Select	02/10/2020	G - No patients consente d	Participant approached within the initiation window but did not consent	Neither

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19/WA/0 325	266292	A Randomised Controlled Trial of Early surgery in severe ASYmptomatic Aortic Stenosis	No		235			10/01/2020	10/01/2020	20/04/2020	26/08/2020	01/09/2020	Please Select	22/09/2020		No participants recruited, still within initiation target window	Please Select
20/NI/000 4	260418	Feasibility Study of the HighLife 28mm Trans-septal Trans-catheter Mitral Valve in Patients With Moderate-Severe or Severe Mitral Regurgitation and at High Surgical Risk	No		235			10/01/2020	10/01/2020	20/04/2020	26/08/2020	01/09/2020	Please Select	22/09/2020		Still within initiation target window	Please Select
19/NE/03 74	275170	MK8591A-017 study – A Phase 3 Randomized, Active-Controlled, Open-Label Clinical Study to Evaluate a Switch to MK- 8591A (Islatravir/Doravirine) Once-Daily in Participants With HIV-1 Virologically Suppressed on Antiretro	No		71			20/12/2020	15/01/2020	17/03/2020	13/03/2020	26/03/2020	Please Select	30/03/2020	J - Other	Recruitment put on hold - COVID 19	Neither
20/EE/01 01	281712	Randomised Evaluation of COVID-19 Therapy (RECOVERY)	Yes	02/04/2020	4	7	11	17/03/2020	22/03/2020	17/03/2020	13/03/2020	26/03/2020	Please Select	30/03/2020		Initiation target met	Please Select
20/SC/01 54	281800	A Multicenter, Adaptive, Randomized Blinded Controlled Trial of the Safety and Efficacy of Investigational Therapeutics for the Treatment of COVID-19 in Hospitalised Adults	Yes	15/04/2020	13	9	22	24/03/2020	24/03/2020	26/03/2020	25/03/2020	06/04/2020	Please Select	06/04/2020		Initiation target met	Please Select
20/YH/00 63	266296	A PHASE 3 OPEN-LABEL, MULTI- CENTER, LONG-TERM STUDY INVESTIGATING THE SAFETY AND EFFICACY OF P-06851600 IN ADULT AND ADOLESCENT PARTICIPANTS WITH ALOPECIA AREATA	No					04/06/2020	13/07/2019				Please Select			Study in set up - not open to recruitment. The sponsor had to go back to REC.	Please Select
18/LO/06 60	237150	Randomized, Embedded, Multifactorial, Adaptive Platform trial for Community- Acquired Pneumonia	Yes	12/06/2020	21	31	52	21/04/2020	21/04/2020	09/01/2020	07/05/2020	12/05/2020	Please Select	26/05/2020		Initiation target met	Please Select
20/SC/02 11	282109	Chloroquine/ hydroxychloroquine prevention of coronavirus disease (COVID-19) in the healthcare setting; a randomised, placebo-controlled prophylaxis study (COPCOV)	Yes	21/05/2020	1	8	9	12/05/2020	12/05/2020	13/05/2020	28/04/2020	13/05/2020	Please Select	21/05/2020		Initiation target met.	Please Select
20/EE/01 35	282213	Multiarm Therapeutic study in pre-ICU patients admitted with COVID-19 - Repurposed Drugs (TACTIC-R)	No		42			09/06/2020	09/06/2020	06/05/2020	21/07/2020	21/07/2020	Please Select	02/08/2020	F - No patients seen	Initiation target not met as COVID pt numbers were low at the time	