Whittington Hospital NHS Trust Performance in Initiating Q2 17/18

Research Ethics Committee Reference Number	Integrated Research Application System Number	Name of Trial	First Patient Recruited?	Date of 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	Benchmark Met	Date Site Invited	Date Site Selected	HRA Approval Date	Date Site Confirmed By Sponsor	Date Site Confirmed	Non-Confirmation Status	Reasons for Delay	Comments
16/WA/0342	215547	Tailored internet-delivered cognitive behavioural therapy for depression and anxiety in patients with a long-term condition (Chronic Pain, COPD and Diabetes)	Yes	01/02/2017	43	14	57	Yes	09/12/2016	06/12/2016	28/11/2016	11/01/2017	18/01/2017	-	-	-
17/LO/0118	220143	Maternal Moments: Investigating music listening for well-being in pregnancy	Yes	11/08/2017	38	67	105	No	17/01/2017	28/04/2017	28/04/2017	28/04/2017	05/06/2017	-	G - No patients consented J - Other	Student had delays with issuing her letter of access
16/WA/0341	209738	Acceptability and Effectiveness of an Internet-Delivered Intervention for Psychological Distress in Patients with Type 2 Diabetes.	Yes	31/07/2017	40	41	81	No	24/04/2017	11/05/2017	18/01/2017	13/06/2017	20/06/2017	-	G - No patients consented	Despite screening no eligible patient has consented to participate in the study. Recruitment has been challenging in all sites, Sponsor is investigating.
17/SC/0244	226900	A randomised, parallel-group, double-blind, double-dummy, active-controlled, multicentre study to assess the efficacy and safety of vilaprisan in subjects with uterine fibroids. ASTEROID 5	Yes	07/09/2017	3	38	41	Yes	15/05/2017	28/07/2017	27/07/2017	31/07/2017	31/07/2017	-	-	-

Whittington Health NHS Trust

Whittington Hospital NHS Trust Performance in Delivery Q2 17/18

Research Ethics Committee Reference Number	Integrated Research Application System Number	Name of Trial	Target Number Of Patients Agreed?	Minimum Number Of Patients Agreed (Enter Same In Both If Only One Number)	Maximum Number Of Patients Agreed (Enter Same In Both If Only One Number)	Target Date To Recruit Patients Agreed?	Date Agreed to recruit target number of patients	Total Number Of Patients Recruited At The Agreed Target Date	Date That The Trial Closed To Recruitment	Total Number Of Study Participants Recruited	Reason For Closure Of Trial	Comments
16/LO/0083	193141	A Phase 3, double-blind, randomized, placebo-controlled, multicenter study to determine the efficacy and safety of luspatercept (ACE-536) plus best supportive care versus best supportive care in adults who require regular red-blood cell transfusions	Number Agreed	3	3	Date Agreed	01/02/2018	2	10/06/2017	2	Withdrawn By Sponsor	The sponsor halted recruitment early, before the target could be reached.