

Performance in Initiating Clinical Research

April 2020 - March 2021

Quarter Reported	Research Ethics Committee Reference Number	Integrated Research Application System Number	Name of Trial	First Participant Recruited?	Date of First Participant Recruited	Date Site Invited	Date Site Selected	HRA Approval Date	Date Site Confirmed By Sponsor	Date Site Confirmed	Date Site Ready To Start	Benchmark Met	Reason For Delay	Comments	Reasons for delay correspond to:
Q3	20/LO/1044	287176	A Randomized, Double-Blind, Placebo-controlled Study Evaluating the Efficacy and Safety of SAGE-217 in the Treatment of Adults with Severe Postpartum Depression	No		18/06/2020	02/09/2020	25/11/2020	31/03/2021	31/03/2021	26/04/2021	No	No patients consented	Contracting delays during set-up due use of non-model commercial agreement. Challenging eligibility criteria required for study.	Both
Q3	19/LO/1667	271363	Clinical and cost-effectiveness of a New psychological intervention to support Independence in Dementia (NIDUS) for family carers and people living with dementia in their own homes: A randomised controlled trial	Yes	27/01/2021	29/06/2020	18/09/2020	13/01/2020	18/09/2020	23/10/2020	10/11/2020	No	Staff availability issues	70 day benchmark missed. Date site selected to confirmed 5 days over. Delays associated with staff availability	NHS Provider
Q4	20/LO/0034	274277	APPLE-Tree programme for dementia prevention: pilot and RCT	Yes	12/07/2021	04/03/2021	25/03/2021	14/01/2020	15/04/2021	19/04/2021	16/06/2021	No	Staff availability issues	70 day benchmark missed but date site selected to confirmed 25 calendar days. Delays were associated with obtaining information required to issue HR arrangements for external researchers and staff availability.	Both