Performance in Initiating Clinical Research Q4 2013-14

70-day benchmark, valid application to first participant recruited

Trial Name	Research Ethics Committee Number	Date of receipt of a Valid Research Application (SSI)	Date of the recruitment of first patient	Reason for not achieving benchmark, if applicable
A pilot study on the effects of adding c-reactive protein point of care testing in acutely ill children in primary care	13/SC/0045	08/07/2013	16/07/2013	
A randomised control trial of guided self-help for binge eating	13/SC/0217	05/08/2013	21/08/2013	
Understanding the importance of plasticity in the brain mechanisms of dyspnoea perception	12/SC/0713	11/07/2013	12/11/2013	Following appropriate governance and granting of NHS Permission, sponsor requested a site agreement, despite no transfer of funds and no additional activity outside routine clinical care.
BCoS-Care: a pilot study for developing and evaluating a care pathway for cognitive problems after stroke	12/WM/0335	08/07/2013	08/09/2013	
An optimised person-centred intervention to improve quality of life for people with dementia living in care homes. A cluster randomised controlled trial	13/SC/0281	08/08/2013	20/11/2013	Permission delayed due to recruitment methodology queries. Managers of care homes needed to agree to their involvement before residents could be approached for recruitment, therefore delaying first patient consent.
OXTEXT-7: Stepped wedge cluster randomised controlled trial of Feeling Well with True Colours for community patients in Oxford Health NHS FT: quantitative evaluation	13/SC/0070	11/06/2013	20/06/2013	
Feasibility study for a community based intervention for individuals with severe CFS/ME	13/SC/0046	05/07/2013	12/03/2014	This study was delayed significantly as Oxford Health NHS FT did not receive all required regulatory documentation that was necessary to begin the study. Once NHS Permission was granted a patient was recruited within 15 days.
Physical activity programmes for community dwelling people with mild to moderate dementia (DAPA- Dementia and physical activity)	11/SW/0232	20/03/2013	24/04/2013	
A randomised, placebo-controlled, parallel-group, double-blind, efficacy and safety trial of MK-8931 in subjects with mild to moderate Alzheimer's Disease	12/NE/0410	08/04/2013	15/05/2013	
A randomised, double-blind, placebo-controlled, phase 3 study to evaluate the efficacy and safety of once a day, TAK-375 (Ramelteon) tablet for sublingual administration (TAK-375SL tablet) 0.1mg and 0.4mg as an adjunctive therapy in the treatment of acute depressive episodes associated with bipolar 1 disorder in adult subjects	12/EM/0391	30/10/2013	20/12/2013	
Mood Action Psychology Programme (MAPP): a case series investigation of brief imagery-focused cognitive therapy (imCT) for Bipolar Disorder	13/EE/0174	08/10/2013	06/11/2013	
Assisted Technology and Telecare to Maintain Independent Living at Home for People with Dementia	12/LO/1816	03/12/2013	12/12/2013	

Trial Name	Research Ethics Committee Number	Date of receipt of a Valid Research Application (SSI)	Date of the recruitment of first patient	Reason for not achieving benchmark, if applicable
Minocycline in Alzheimer's Disease efficacy trial: The MADE Trial	13/EE/0063	13/03/2014		Patient not yet recruited. The 70 day benchmark is 21/05/2014
Long-term safety and efficacy of ABT-126 in subjects with schizophrenia: a double-blind extension study for subjects completing M10-855	13/SC/0386	22/01/2014		Patient not yet recruited. The 70 day benchmark is 02/04/2014
A randomised, double-blind, placebo-controlled, single-dose study of the effects of SEP 363856 and Amisulpride on bold-FMRI signal in healthy male and female volunteers with high or low schizotype characteristics	13/NW/0727	07/03/2014	27/03/2014	