

Performance in Initiating Clinical Trials Q2 2016-2017  
70 day benchmark, valid application to first participant recruited  
NHS Permission Studies only

| Research Ethics Committee Number | IRAS Number | Name of Trial  | Receipt of Valid Research Application | Date of First Patient Recruited | Benchmark Met | Comments  |
|----------------------------------|-------------|--|---------------------------------------|---------------------------------|---------------|---|
| 15/SC/0502                       | 184517      | The Nightmare Intervention Study: a pilot randomised controlled trial of a brief cognitive behavioral therapy for nightmares for patients with persecutory delusions   | 23/11/2015                            | 01/02/2016                      | Yes           |   |
| 15/SS/0032                       | 165287      | STARBEAM Ext: An open-label extension study to evaluate the long-term safety and tolerability of Lu AE58054 as adjunctive treatment to donepezil in patients with mild-moderate Alzheimer's disease  | 09/02/2016                            | 01/03/2016                      | Yes           |   |
| 14/LO/2071                       | 160786      | Preventing enduring behavioural problems in young children through early psychological intervention: Healthy Start, Happy Start  | 08/01/2016                            | 12/02/2016                      | Yes           |   |
| 15/SC/0508                       | 186095      | Psychological support for fears about other people: A comparison of the Feeling Safe Programme to befriending  | 22/01/2016                            | 08/02/2016                      | Yes           |   |
| 15/YH/0051                       | 172273      | A Smoking Cessation Intervention for severe Mental Ill Health Trial (SCIMITAR+) : a definitive randomised evaluation of a bespoke smoking cessation service  | 29/03/2016                            | 17/05/2016                      | Yes           |   |
| 15/SC/0432                       | 181387      | A Randomized, Double-blind, Multicenter, Active-controlled Study to Evaluate the Efficacy, Safety, and Tolerability of Intranasal Esketamine Plus an Oral Antidepressant in Elderly Subjects with Treatment-resistant Depression - TRANSFORM | 28/03/2016                            |                                 | No            | The sponsor recognised study is difficult to recruit to for all sites and has extended recruitment and introduced a new recruitment tool. |
| 15/SC/0434                       | 181955      | An open-label, long-term, safety and efficacy study of intranasal Esketamine in Treatment-resistant Depression in Adults SUSTAIN-2   | 28/03/2016                            | 17/06/2016                      | No            | This study has been difficult to recruit to for all sites. Sponsor has initiated a new recruitment tool.                                  |
| 15/NE/0269                       | 188082      | A Parallel-Group, Double Blind Long Term Safety Trial of MK-8931 in Subjects with Alzheimer's Disease. (Protocol No. MK-8931-017-12) - EPOCH Ext   | 05/04/2016                            | 04/07/2016                      | No            | A sponsor error meant that the study site was not able to start recruitment despite being ready.  |