Performance in Delivering Clinical Research

Time to Target (Commercial Contracts)

Trial Name	Research Ethics Committee Number	Target number of patients	Recruitment end date	Trial Status	Target met?	Comments
RO4917838 in stable patients with persistent, predominant negative symptoms of schizophrenia treated with antipsychotics - Protocol No. NN25310	10/H0606/69	6	08/05/2013	Closed – in follow up	No	Four participants recruited from six, viewed as successful by the sponsor because this study has proven difficult to recruit to across all UK sites. In addition the projected recruitment target for this site was probably over ambitious.
RO4917838 in patients with sub-optimally controlled symptoms of schizophrenia treated with antipsychotics. Protocol No. NN25307	10/H0604/94	6	30/06/2013	Closed – in follow up	No	Again this study has proven difficult to recruit to across all UK sites. In addition the projected recruitment target for this site was probably over ambitious.
Evaluation of the effects of agomelatine (25mg and 50mg) and escitalopram 20mg during 8 weeks on emotional blunting, emotional processing and motivation in healthy male and female volunteers.	11/SC/0383	64	15/05/2013	Closed – in follow up	Yes	
A randomised, double-blind, parallel-group, active-controlled, flexible dose study evaluating the effects of LuAA21004 versus agomelatine in adult patients suffering from major depressive disorder with inadequate response to antidepressant treatment. (REVIVE)	11/NW/0747	8	24/08/2012	Withdrawn	N/A	Recruitment halted by sponsor as target met at other sites that opened before this site. OHFT is a site relying entirely on referrals from a PIC. Of the few referrals made all failed screening. NHS permission delayed due to contractual issues, which reduced time for recruitment.
Interventional randomised, double-blind, parallel-group, placebo-controlled, exploratory study investigating the effects of LuAA21004 on cognition and BOLD fMRI signals in subjects remitted from depression and controls	12/SC/0213	32	31/07/2013	Open	N/A	The study is still open; but expected to complete on target.
A double-blind, Placebo-controlled, Randomised, 4-week safety and Tolerability Study of LMTM in subjects with Mild to Moderate Alzheimer's Disease on Pre-Existing stable Acetylcholinesterase Inhibitor and/or Memantine Therapy	12/NW/0441	4	31/12/2012	Withdrawn	N/A	Sponsor stopped the study early due to lack of recruitment across the country and newer studies using the same IMP were initiated.
A randomised, double-blind, placebo-controlled, dose-ranging, parallel-group, phase 2 study of the safety and efficacy of ABT-126 in the treatment of cognitive deficits in schizophrenia (CDS) in non-smokers	12/SC/0544	3	30/09/2013	Open	N/A	Recruitment started and is expected to complete on target
A randomised, placebo-controlled, parallel-group, double-blind, efficacy and safety trial of MK-8931 in subjects with mild to moderate Alzheimer's Disease (EPOCH)	12/NE/0410	8	31/08/2015	Open	N/A	Recruitment started.