South London and Maudsley NHS Foundation Trust: Q4 2015-2016 Performance in Delivering Research

ld	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)		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	Reason For Closure Of Trial	Comments
			Observational Study Description									
			F1D-MC-B034(a) Post-Injection Syndrome in	Danas			Not Available /				\	
2939	10/47015/26	51787		Range Agreed	11	11			0	24/07/2015	Withdrawn	0 Patient
2939	10/H7015/26	31707	Olanzapine Long-Acting Injection Protocol H8A-MC-LZAO Continued Efficacy	Agreed	11	11	Not Agreed		U	31/07/2015	By Host	U Patient
			and Safety Monitoring of Solanezumab, an				Not					
				Range			Available /				Recruitment	
2940	10/H1102/85	66298		Agreed	1	5	Not Agreed		5	26/10/2015	Finished	5 patients
2540	10/111102/00	00230	Randomized, Double-blind,	rigicca	'		140t Agreed		0	20/10/2010	Tillioned	o patiento
00.44	40/10/4400	105074	Placebo-controlled, Cross-over Study to Evaluate Functional Magnetic Resonance Imaging (fMRI) of the Brain with Administration of Risperidone, Olanzapine, Haloperidol, Alprazolam or Placebo in Healthy		40	40	Data Assard	04/00/0045		00/44/0044	Recruitment	CO Patier
2941	13/LO/1183	135274	Subjects	Agreed	42	42	Date Agreed	31/03/2015	63	03/11/2014	Finished	63 Patien
2942	13/LO/0702	116726	A Randomized, Double-blind, Placebo-controlled, Parallel, 26-Week, Phase 3 Study of 2 Doses of an Alpha-7 Nicotinic Acetylcholine Receptor Agonist (EVP-6124) or Placebo as an Adjunctive Pro-cognitive Treatment in Schizophrenia Subjects on Chronic Stable At A Randomized, Double-Blind, Placebo	Range Agreed	1	6	Date Agreed	31/10/2014	5	21/05/2015	Recruitment Finished	7 Patient
00.40	10/00/0440		Controlled, 3-period, Proof of Mechanism, Cross-Over Study of Roflumilast (100 or 250 ?g) administered up to Steady State to Evaluate the Effects of Add-on Roflumilast to Second Generation Antipsychotics on	Range		4-5		0.1/10/001.1		05/00/0045	Recruitment	
2943	12/SC/0443	111575	Cognitive Imp A Multicenter 26-Week Extension Study to	Agreed	1	15	Date Agreed	31/12/2014	4	25/09/2015	Finished	5 patients
2944	14/LO/0359	131838	Evaluate the Safety and Clinical Effects of Prolonged Exposure to 1 and 2 mg Doses of EVP-6124, an Alpha- 7 Nicotinic Acetylcholine Receptor Agonist, as an Adjunctive Pro-cognitive	Range Agreed	1	6	Date Agreed	31/12/2015	4	14/09/2015	Withdrawn By Sponsor	4 Patient