Research Ethics Committee Reference Number	Full Name of Trial	Target number of patients	Date Agreed to recruit target number of patients	Trial Status	Target met within the agreed time
11/SC/0524	Phase 3b Randomized Open Label Study to Evaluate Switching from Regimens Consisting of a NNRTI plus Emtricitabine (FTC) and TDF to EVG/COBI/FTC/TDF in VirologicallySuppressed, HIV1 Infected Patients	5	31/12/2012	Closed - Follow Up Complete	Y
12/SC/0035	A 6-month safety and benefit study of inhaled fluticasone propionate/ salmeterol combination versus inhaled fluticasone propionate in the treatment of 6,200 paediatric subjects 4-11 years old with persistent asthma	2	31/08/2015	Closed - Follow Up Complete	Y
12/EE/0176	Randomisd Ph 4 placebo-controlld comparative study to evaluate efficacy/safety of tapering MTX dosage vs maintaining dosage in severe active RA patients showing inadeq response to prior conventional DMARDs trtmt & initiatd RoActemra? in combo w/ MTX	5	01/04/2014	Closed - Follow Up Complete	Y
11/LO/1921	Randomized 2 arm open-label multicenter Phase 2 trial assessing efficacy & safety of pertuzumab given in combo w/trastuzumab & aromatase inhibitor in 1st line pts w/HER2+ & hormone receptor+ advanced (metastatic/locally advanced) breast cancer	5	29/01/2016	Closed - Follow Up Complete	N
10/H0808/137	GORE HELEX Septal Occluder and Antiplatelet Medical Management for Reduction of Recurrent stroke or Imaging-confirmed TIA in patients with Patent Foramen Ovale (PFO)	20	04/12/2017	Closed - In Follow Up	N

12/NI/0181	REPRISE II: REpositionable Percutaneous Replacement of Stenotic Aortic Valve through Implantation of Lotus? Valve System ? Evaluation of Safety and Performance.	6	31/01/2014	Closed - in follow up	Y
11/SC/0329	Phase 3 Randomized Double-blind Placebo-controlled Parallel-group Multicenter Study to Evaluate Safety/Efficacy of Ustekinumab Maintenance Therapy in Subjects w/ Mod/Severe Active Crohn?s Disease Who Failed/Are Intolerant to TNF Antagonist Therapy	1	01/10/2017	Closed - In Follow Up	Y
11/LO/1498	PROTEAse inhibitor (DRV/rtv) in mono- or triple therapy in suppressed HIV-1 infected subjects.	10	31/01/2013	Closed - Follow Up Complete	Υ
10/H1102/85	Protocol H8A-MC-LZAO Continued Efficacy and Safety Monitoring of Solanezumab, an Anti- Amyloid ? Antibody in Patients with Alzheimer?s Disease	N/A	N/A	Closed - In Follow Up	Υ
11/LO/0537	A Multicenter, Randomized, Double-Blind, Placebo-Controlled Study of the Efficacy of Natalizumab on Reducing Disability Progression in Subjects With Secondary Progressive Multiple Sclerosis	6	27/02/2015	Closed - In Follow Up	Y
12/NW/0723	A multi-centre, open-label, long term safety extension of phase II studies ABE4869g and ABE4955g in patients with mild to moderate Alzheimer's Disease.	4	N/A	Closed - In Follow Up	Υ

11/LO/1381	A phase II double-blind placebo-controlled randomized study of GDC-0941 or GDC-0980 with Fulvestrant versus Fulvestrant in advanced or metastatic breast cancer in patients resistant to aromatase inhibitortherapy	5	31/01/2013	Closed - Follow Up Complete	Y
13/LO/0033	Genentech GO28509-PEGGY: A PHASE II, randomized STUDY OF paclitaxel with GDC- 0941 versus paclitaxel with placebo IN PATIENTS WITH LOCALLY RECURRENT OR METASTATIC BREAST CANCER	3	01/09/2014	Closed - Follow Up Complete	Y
12/SW/0378	Effect of Bivalirudin on Aortic Valve Intervention outcomes 2/3	20		Closed - Follow Up Complete	N
13/LO/0574	Phase 3 Randomized Double-Blind Study to Evaluate Safety/Efficacy of Elvitegravir/Cobicistat/Emtricitabine/ Tenofovir Alafenamide vs Elvitegravir/Cobicistat/ Emtricitabine/Tenofovir Disoproxil Fumarate in HIV 1+ Antiretroviral Treatment-Naive Adults	10	23/07/2013	Closed - In Follow Up	N
13/LO/0821	A Phase 3 Open-label Safety Study of Elvitegravir/Cobicistat/Emtricitabine/Tenofovir Alafenamide Single-Tablet Regimen in HIV-1 Positive Patients with Mild to Moderate Renal Impairment	5	31/12/2013	Closed - In Follow Up	N
13/LO/0830	Phase 3 open label study evaluating efficacy/safety of pegylated interferon lambda-1a, in combination + ribavirin and daclatasvir, for treatment of chronic HCV infection + treatment na?ve genotypes 1, 2, 3 or 4 in subjects co-infected + HIV	5	31/10/2014	Closed - Follow Up Complete	N

13/NW/0002	Prospective, randomised, controlled investigation comparing the safety and performance of 032-11 Surgical Haemostat 2g Applicator with FLOSEAL? Haemostatic Matrix as an adjunctive haemostat in cardiac surgery and thoracic aortic surgery	10	28/02/2014	Open	Y
13/NW/0316	Phase 4 trial comparing cumulative incidence of SCC after treatment + ingenol mebutate & imiquimod for multiple actinic keratoses on face &scalp. A multi-centre, randomised, two-arm, open label, active-controlled, parallel group, 36-month trial.	8	31/05/2015	Open	Y
13/SC/0279	A Phase 3, Open-Label Study to Evaluate Switching from a TDF-Containing Combination Regimen to a TAF-Containing Combination Single Tablet Regimen (STR) in Virologically Suppressed, HIV1 Positive Subjects.	Rollover from GS 0106/0114/010 3 no target	17/01/2014	Closed - In Follow Up	N/A
11/NW/0597	An Open-Label, Dose-Escalation, Phase 1/2 Study of the Oral Form of MLN9708, a Next- Generation Proteasome Inhibitor, Administered in Combination With a Standard Care Regimen of Melphalan and Prednisone in Patients With Newly Diagnosed Multiple Myeloma Req	2	30/06/2015	Closed - In Follow Up	N
13/NW/0501	A Phase 3, Randomized, Double-blind, Placebo-controlled Study to compare efficacy and safety of Oral Azacitidine plus best supportive care versus best supportive care as Maintenance Therapy in subjects with Acute Myelogenous LeukEmia in complete remission	3	15/02/2018	Open	Y
13/LO/0671	A Multicenter, Open-Label, Extension Study to Evaluate the Long-Term Safety and Efficacy of BIIB019, Daclizumab High Yield Process (DAC HYP), Monotherapy in Subjects With Multiple Sclerosis Who Have Completed Study 205MS301.	3	05/02/2014	Closed - In Follow Up	Y

13/NW/0560	Effect of Passive Immunization on the Progression of Mild Alzheimer's Disease: Solanezumab (LY2062430) Versus Placebo	8	20/06/2014	Closed - In Follow Up	Ν
13/LO/0908	A Phase 2, Single-Arm, Open-Label, Multicenter Study of the Clinical Activity and Safety of Enzalutamide in Patients With Advanced, Androgen Receptor-Positive, Triple-Negative Breast Cancer	2	30/12/2015	Closed - In Follow Up	Y
13/LO/1720	A Phase 2, Randomized, Double Blind, Placebo Controlled, Multicenter Study of Efficacy and Safety of Enzalutamide in Combination With Exemestane in Patients With Advanced Breast Cancer That Is Estrogen or Progesterone Receptor Positive and HER2 Normal	6	15/04/2016	Closed - Follow Up Complete	N
14/SC/0225	Gilead 311-1089 Phase 3 randomised open-label switch study to evaluate F/TAF in HIV-1 positive subjects who are virologically suppressed on regimens containing FTC/TAF	8	31/05/2016	Closed - in follow up	Ν
13/YH/0282	ACT-MOVE: ML28641 - Subcutaneous tocilizumab in rheumatoid arthritis	3	30/06/2015	Closed - In Follow Up	Y
13/EE/0126	Evaluation of Safety and Efficacy of the BACE™ [Basal Annuloplasty of the Cardia Externally] Device in the Treatment of Functional Mitral Valve Regurgitation [FMR]	6	01/07/2016	Closed - Follow Up Complete	N
14/LO/0081	A Phase IIb, Multi-Center, Randomized, Double-Blind, Placebo-Controlled, Multidose, 24- Week Study to Evaluate the Efficicacy and Safety of Atacicept in Subjects With Systemic Lupus Erythematosus	3	31/07/2015	Closed - Follow Up Complete	N

14/NW/0017	A Prospective, Single-Arm, Clinical-Setting Study to Describe Efficacy, Tolerability and Convenience of Teriflunomide Treatment Using Patient Reported Outcomes (PROs) in Relapsing Multiple Sclerosis (RMS) Patients.	6	30/11/2015	Closed - in follow up	Y
13/LO/1081	PROSPER: A Multinational, Phase 3, Randomized, Double-Blind, Placebo-Controlled, Efficacy and Safety Study of Enzalutamide in Patients With Nonmetastatic Castration- Resistant Prostate Cancer	4	01/02/2016	Closed - Follow Up Complete	N
13/NE/0126	An exploratory, randomised, double-blind, controlled study to assess the effect of an Amino Acid Based Formula with a synbiotic blend on gut microbiota and stool characteristics in infants with suspected gastrointestinal Non IgE mediated cow's milk allergy (CMA).	5	28/07/2014	Closed - Follow Up Complete	Y
14/SW/0079	A Prospective, Randomized Evaluation of the TriGuard™ HDH Embolic DEFLECTion Device during Transcatheter Aortic Valve Implantation	6	01/12/2014	Closed - Follow Up Complete	N
14/EM/0129	A PHASE 2/3, MULTI-CENTRE, RANDOMISED, DOUBLE-BLIND, PLACEBO- CONTROLLED (PART A) AND DOUBLE-BLIND, DOUBLE-DUMMY, ACTIVE- CONTROLLED (PART B), PARALLEL GROUP STUDY TO EVALUATE THE EFFICACY AND SAFETY OF RPC1063 ADMINISTERED ORALLY TO RELAPSING MULTIPLE SCLEROSIS PATIENTS	6	01/05/2015	Closed - In Follow Up	N
14/WS/0004	Open-Label Extension Study of EFC12492, R727-CL-1112, EFC12732, & LTS11717 Studies to Assess the Long-Term Safety and Efficacy of Alirocumab in Patients with Heterozygous Familial Hypercholesterolmia	2	31/10/2014	Closed - In Follow Up	Y

14/LO/1381	Reformulated raltegravir q.d. (1200 mg) versus raltegravir b.i.d. (400 mg) in ART-naïve pts	5	30/04/2015	Closed - In follow up	Y
14/LO/1513	A Randomized, Open Label, Phase 4 Study Evaluating the Renal Effect of Elvitegravir/Cobicistat/Emtricitabine/Tenofovir DF or other Tenofovir DF-containing Regimens (Ritonavir-boosted Atazanavir plus Emtricitabine/Tenofovir DF or Efavirenz /Emtricitabine/Tenofovir DF) compared to Ritonavir boosted Atazanavir plus Abacavir/Lamivudine in Antiretroviral Treatment-naïve HIV-1 Infected Adults with eGFR ≥70 mL/min	3	31/12/2015	Closed - in follow up	Y
14/LO/0298	A multicenter, Single Arm Study of Enzalutamide in Patients with Progressive Metastatic Castration-Resistant Prostate Cancer Previously Treated With Abiraterone Acetate.	10	01/08/2016	Closed - In Follow Up	N
14/LO/1052	A Phase 2, Randomized, OpenLabel,Parallel Group Study Evaluating the Safety and Efficacy of TAK385, an Oral Gonadotropin Releasing Hormone (GnRH) Antagonist, for Patients With Localized Prostate Cancer Requiring Neoadjuvant and Adjuvant Androgen Deprivation Therapy With External Beam Radiation Therapy (EBRT)	2	N/A	Closed - In Follow Up	Y
14/YH/0088	Phase I study of KHK2823 in Patients with Acute Myeloid Leukaemia or Myelodysplastic Syndrome	4	01/12/2017	Open	N/A
14/YH/0086	RESPOND: Repositionable Lotus Valve System-Post Market Evaluation of Real World Clincial Outcomes	15	31/12/2015	Open	Y

14/SC/1161	Prospective, single-arm, Multi-centre, observational registry to Further Validate Safety and Efficacy of the Ultimaster DES system in unselected patients representing everyday clinical practice	30	01/12/2015	Open	Y
13/LO/1891	More Response on Cardiac Resynchronization Therapy (CRT) with MultiPoint Pacing (MPP)	15	01/12/2016	Open	Y
13/EM/0348	Safety and Efficacy assessment of Monoprost® (unpreserved latanoprost) in comparison with Lumigan® 0.01 % and Lumigan® 0.03% UD, in patients with primary open angle glaucoma or ocular hypertension, stabilized by Lumigan® 0.01 % with ocular surface intoler	6	30/04/2016	Open	N/A
13/YH/0147	A Randomized, Open Label, Multi-Centre, Controlled Study to Assess Safety and Efficacy of ELAD in Subjects with Acute Alcoholic Hepatitis (AAH) Who Have Failed Steroid Therapy (incorporating VTI-210E as a follow-up registry)	4	31/08/2015	Closed - Follow Up Complete	N
14/WM/0013	A Randomized, Double blind, Placebo-Controlled, 2-Part Study of Orally Administered ALS- 008176 to Evaluate the Safety, Tolerability, Pharmacokinetics and Pharmacodynamics of Single Ascending Dosing and Multiple Ascending Dosing in Infants Hospitalized with Respiratory Syncytial Virus (RSV) Infection.	4	30/04/2015	Open	N
14/EM/1070	A Phase III, Randomized, Double-Blind, Placebo-Controlled, Multicenter Clinical Study Evaluating the Safety and Efficacy of Icatibant as a Treatment for Angiotensin-Converting Enzyme Inhibitor (ACE-I)-Induced Angioedema in Adults	1	01/10/2015	Closed - in follow up	Y

14/LO/1443	A Phase 3, Randomized, Open-Label, Active-Controlled Study to Evaluate the Efficacy and Safety of FG-4592 in the Treatment of Anemia in End Stage Renal Disease Subjects on Stable Dialysis Converted from Epoetin or Darbepoetin Alfa Treatment	4	19/12/2017	Open	N/A
14/LO/1435	A Double-Blind, Randomized, Parallel-Group, Active-Control Study to Compare the Efficacy and Safety of CHS-0214 Versus Enbrel® in Subjects With Rheumatoid Arthritis and Inadequate Response to Treatment With Methotrexate	3	30/05/2016	Closed - Follow Up Complete	N
13/YH/0315	A randomized, parallel group, open-label, multicentre study to investigate the efficacy and safety of oral BAY 85-3934 and active comparator (darbepoetin alfa) in the maintenance treatment of anemia in pre-dialysis subjects with chronic kidney disease on and Asia Pacificdarbepoetin treatment in Europe	1	31/03/2015	Closed - In Follow Up	N
14/LO/0892	A human immunoglobulin G1 kappa (IgG1κ) monoclonal antibody directed against programmed death ligand 1 (PD-L1)	3	15/02/2015	Open	N
14/SS/1048	A Phase 3b, Multi-center, Randomized-withdrawal, Placebo-controlled, Double blind, Parallel-group Trial to Compare the Efficacy and Safety of Tolvaptan (45 to 120 mg/day, Split-dose) in Subjects with Chronic Kidney Disease Between Late Stage 2 to Early Stage 4 Due to Autosomal Dominant Polycystic Kidney Disease	5	10/07/2015	Open	N
13/NI/0188	A single-arm trial to evaluate the effectiveness of PCI of de novo 3-vessel disease applying the SYNTAX Score II with pressure wire functional assessment and IVUS guidance, using an everolimus-eluting stent with biodegradable abluminal coating	10		Closed - In Follow Up	N

	A Desce 2 rendemized active controlled once level study to evolve to the efficiency of the				
	A Phase 3, randomized, active-controlled, open-label study to evaluate the efficacy, safety				
	and tolerability of switching to a darunavir/cobicistat/emtricitabine/tenofovir alafenamide			0	
	(D/C/F/TAF) once-daily single-tablet regimen versus continuing the current regimen			Open	
4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4	consisting of a boosted protease inhibitor (bPI) combined with emtricitabine/tenofovir		42/00/2015		V
14/WM/1210	disoproxil fumarate (FTC/TDF) in virologically-suppressed, human immunodeficiency	4	12/08/2015		Y
44/007	A Phase 3b, Multi-center, Open-label Trial to Evaluate the Long Term Safety of Titrated	45	02/00/2015		
44/SS/1087	Immediate-release Tolvaptan (OPC 41061, 30 mg to 120 mg/day, Split dose) in Subjects	15	02/09/2015	Open	N
	with Autosomal Dominant Polycystic Kidney Disease				
	Effects of ODM-109 on respiratory function in patients with ALS. A randomised, double	-	42/40/2045	0	N
15/LO/0684	blind, placebo-controlled, cross-over, 3-period, multicentre study with open-label follow-	5	13/10/2015	Open	Ν
	up extension				
	A Phase 3 Multicenter, Double-Blind, Randomized, Active Comparator-Controlled Clinical				
	Trial to Evaluate the Safety and Efficacy of Doravirine (MK-1439) 100 mg Once Daily Versus	-	01/03/2018	Closed - In	
15/LO/0075	Darunavir 800 mg Once Daily plus Ritonavir 100 mg Once Daily, Each in Combination with	S	01/03/2018	Follow Up	Ν
	TRUVADA™ or EPZICOM™/KIVEXA™, in Treatment-Naïve HIV-1 Infected Subjects				
		4 Healthy			
		Volunteers 10			
		HIV, 4			
	Comparative Testing of 3 mL TransFix/EDTA Vacumm Blood Collection Tubes (TVTs) and	Leukaemia			
14/NW/1531	Cyto-Chex 5 mL Blood Collection Tubes (BCTs) Part 1: Equivalence Study	Patients	05/11/2015	Open	Y
15/YH/0045	ABLATOR Ablation Observational Registry	25	17/07/2017	-	N/A
13/NI/0138	Portico I Study International long-term follow-up studyt of patients	10	30/06/2018	Open	N/A
	COOL-AMI EU CASE SERIES CLINICAL STUDY: a single-centre case series clinical study to				
	assess the feasibility of integrating therapeutic hypothermia(TH)using the ZOLL IVTM				
	System as an adjuvant therapy in percutaneous coronary intervention (PCI) in patients with				
13/EE/0335	acute myocardial infarction (AMI)	5	28/02/2016	Open	N/A
	Randomised Evaluation of dabigatran etexilate Compared to warfarIn in pulmonaRy vein				
	ablation: assessment of an uninterrupted periproCedUral antIcoagulation sTrategy (The RE-				
15/SC/0280	CIRCUIT Trial)	9	31/08/2016	Open	N/A
	Open-label evaluation of the population pharmacokinetic profile, safety, tolerability, and				
	efficacy of intravenous tapentadol solution for injection for the treatment of post-surgical				
14/YH/1269		2	21/00/2016	Onon	
14/10/1209	pain in children aged from birth to less than 2 years, including preterm neonates	3	31/08/2016	Open	N/A

A Phase III Multicenter, Double Blind, Randomized, Active Comparator-Controlled Clinical Trial to Evluate the Safety and Efficacy of MK-149A One Daily Versus ATRIPLA Once-Daily in Treatment Naïve HIV - 1 Infected Subjects 5 28/02/2   A Phase III Multicenter, Open-Label, Randomized Study to Evaluate a Switch to MK-1439A in HIV-1-Infected Subjects Virologically Suppressed on a Regimen of a Ritonavir-boosted 5 19/02/2   15/NW/0505 Protease Inhibitor and Two Nucleoside Reverse Transcriptase Inhibitors (NRTIs) 5 19/02/2   15/LO/1324 AMPLATZER Amulet Observational Post-Market Study 10 17/11/2   A Phase 3, randomized, active-controlled, double-blind study to evaluate efficacy and safety of darunavir/cobicistat/emtricitabine/tenofovir alafenamide (D/C/F/TAF) once daily fixed dose combination regimen versus a regimen consisting of darunavir/cobicistat fixed dose combination coadministered with emtricitabine/tenofovir disoproxil fumarate fixed dose combination in antiretroviral treatment-naïve human immunodeficiency virus type 1		
A Phase III Multicenter, Double Blind, Randomized, Active Comparator-Controlled Clinical Trial to Evluate the Safety and Efficacy of MK-149A One Daily Versus ATRIPLA Once-Daily in 5/LO/0881 5   Treatment Naïve HIV - 1 Infected Subjects 5   A Phase III Multicenter, Open-Label, Randomized Study to Evaluate a Switch to MK-1439A in HIV-1-Infected Subjects Virologically Suppressed on a Regimen of a Ritonavir-boosted 5   5/LO/0505 Protease Inhibitor and Two Nucleoside Reverse Transcriptase Inhibitors (NRTIs) 5 19/02/2   5/LO/1324 AMPLATZER Amulet Observational Post-Market Study 10 17/11/2   A Phase 3, randomized, active-controlled, double-blind study to evaluate efficacy and safety of darunavir/cobicistat/emtricitabine/tenofovir alafenamide (D/C/F/TAF) once daily fixed dose combination regimen versus a regimen consisting of darunavir/cobicistat fixed dose combination coadministered with emtricitabine/tenofovir disoproxil fumarate fixed dose combination in antiretroviral treatment-naïve human immunodeficiency virus type 1 5   5/LO/1239 infected subjects. 5 30/09/2		
Trial to Evluate the Safety and Efficacy of MK-149A One Daily Versus ATRIPLA Once-Daily in Treatment Naïve HIV - 1 Infected Subjects515/L0/0881Treatment Naïve HIV - 1 Infected Subjects5A Phase III Multicenter, Open-Label, Randomized Study to Evaluate a Switch to MK-1439A in HIV-1-Infected Subjects Virologically Suppressed on a Regimen of a Ritonavir-boosted19/02/215/L0/1324AMPLATZER Amulet Observational Post-Market Study1017/11/2A Phase 3, randomized, active-controlled, double-blind study to evaluate efficacy and safety of darunavir/cobicistat/emtricitabine/tenofovir alafenamide (D/C/F/TAF) once daily fixed dose combination regimen versus a regimen consisting of darunavir/cobicistat fixed dose combination in antiretroviral treatment-naïve human immunodeficiency virus type 1515/L0/1239infected subjects.530/09/2	15 Open	Ν
15/L0/0881 Treatment Naïve HIV - 1 Infected Subjects 5 28/02/2   A Phase III Multicenter, Open-Label, Randomized Study to Evaluate a Switch to MK-1439A 10 19/02/2   In HIV-1-Infected Subjects Virologically Suppressed on a Regimen of a Ritonavir-boosted 5 19/02/2   15/L0/1324 AMPLATZER Amulet Observational Post-Market Study 10 17/11/2   A Phase 3, randomized, active-controlled, double-blind study to evaluate efficacy and safety of darunavir/cobicistat/emtricitabine/tenofovir alafenamide (D/C/F/TAF) once daily fixed dose combination regimen versus a regimen consisting of darunavir/cobicistat fixed dose combination in antiretroviral treatment-naïve human immunodeficiency virus type 1 5 30/09/2   15/L0/1239 infected subjects. 5 30/09/2		
A Phase III Multicenter, Open-Label, Randomized Study to Evaluate a Switch to MK-1439A in HIV-1-Infected Subjects Virologically Suppressed on a Regimen of a Ritonavir-boosted   15/NW/0505 Protease Inhibitor and Two Nucleoside Reverse Transcriptase Inhibitors (NRTIs) 5 19/02/2   15/LO/1324 AMPLATZER Amulet Observational Post-Market Study 10 17/11/2   A Phase 3, randomized, active-controlled, double-blind study to evaluate efficacy and safety of darunavir/cobicistat/emtricitabine/tenofovir alafenamide (D/C/F/TAF) once daily fixed dose combination regimen versus a regimen consisting of darunavir/cobicistat fixed dose combination coadministered with emtricitabine/tenofovir disoproxil fumarate fixed dose combination in antiretroviral treatment-naïve human immunodeficiency virus type 1 5 30/09/2   15/LO/1239 Infected subjects. 5 30/09/2		
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15/NW/0505 Protease Inhibitor and Two Nucleoside Reverse Transcriptase Inhibitors (NRTIs) 5 19/02/2   15/LO/1324 AMPLATZER Amulet Observational Post-Market Study 10 17/11/2   A Phase 3, randomized, active-controlled, double-blind study to evaluate efficacy and safety of darunavir/cobicistat/emtricitabine/tenofovir alafenamide (D/C/F/TAF) once daily fixed dose combination regimen versus a regimen consisting of darunavir/cobicistat fixed dose combination coadministered with emtricitabine/tenofovir disoproxil fumarate fixed dose combination in antiretroviral treatment-naïve human immunodeficiency virus type 1 5 30/09/2   15/LO/1239 infected subjects. 5 30/09/2		
15/LO/1324 AMPLATZER Amulet Observational Post-Market Study 10 17/11/2   A Phase 3, randomized, active-controlled, double-blind study to evaluate efficacy and safety of darunavir/cobicistat/emtricitabine/tenofovir alafenamide (D/C/F/TAF) once daily fixed dose combination regimen versus a regimen consisting of darunavir/cobicistat fixed dose combination coadministered with emtricitabine/tenofovir disoproxil fumarate fixed dose combination in antiretroviral treatment-naïve human immunodeficiency virus type 1 5 30/09/2		
A Phase 3, randomized, active-controlled, double-blind study to evaluate efficacy and safety of darunavir/cobicistat/emtricitabine/tenofovir alafenamide (D/C/F/TAF) once daily fixed dose combination regimen versus a regimen consisting of darunavir/cobicistat fixed dose combination coadministered with emtricitabine/tenofovir disoproxil fumarate fixed dose combination in antiretroviral treatment-naïve human immunodeficiency virus type 1 15/LO/1239 infected subjects. 5 30/09/2	18 Open	N/A
safety of darunavir/cobicistat/emtricitabine/tenofovir alafenamide (D/C/F/TAF) once daily fixed dose combination regimen versus a regimen consisting of darunavir/cobicistat fixed dose combination coadministered with emtricitabine/tenofovir disoproxil fumarate fixed dose combination in antiretroviral treatment-naïve human immunodeficiency virus type 1 15/LO/1239 infected subjects. 5 30/09/2	19 Open	N/A
safety of darunavir/cobicistat/emtricitabine/tenofovir alafenamide (D/C/F/TAF) once daily fixed dose combination regimen versus a regimen consisting of darunavir/cobicistat fixed dose combination coadministered with emtricitabine/tenofovir disoproxil fumarate fixed dose combination in antiretroviral treatment-naïve human immunodeficiency virus type 1 15/LO/1239 infected subjects. 5 30/09/2		
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dose combination coadministered with emtricitabine/tenofovir disoproxil fumarate fixed dose combination in antiretroviral treatment-naïve human immunodeficiency virus type 115/LO/1239infected subjects.530/09/2		
dose combination in antiretroviral treatment-naïve human immunodeficiency virus type 1 15/LO/1239 infected subjects. 5 30/09/2		
15/LO/1239   infected subjects.   5   30/09/2		
Safety and Performance Registry for an all-comers patient population with the Linus	15 OPEN	N
	16 Open	N/A
A Multicentre, Randomised, Double-blind, Placebo-controlled, Phase 3 Study Evaluating the		
Efficacy and Safety of Two Doses of Anifrolumab in Adult Subjects with Active Systemic		
	17 OPEN	N/A