Study title	Name of sponsor	Location for study	Commencement of administrative procedure
Inspection not focused on a single clinical trial	-	MEDICAL PLUS, s.r.o	No
A First in Human, Open-Label Phase I/Ib study to evaluate the Safety, Tolerability, Pharmacokinetics and Clinical Activity of MitoTam in patients with advanced solid tumors	Smart Brain s.r.o.	Všeobecná fakultní nemocnice v Praze	No
Efficacy and safety of low doses of trazodone in patients affected by painful diabetic neuropathy: randomized, controlled, pilot study.	Angelini S.p.A.	Vestra Clinics s.r.o.	No
A First in Human, Open-Label Phase I/Ib study to evaluate the Safety, Tolerability, Pharmacokinetics and Clinical Activity of MitoTam in patients with advanced solid tumors	Smart Brain s.r.o.	Smart Brain s.r.o. - sponsor site inspection	No
A 2-year Prospective Study to Assess Health-related Quality of Life in Subjects with Highly-Active Relapsing Multiple Sclerosis Treated with Mavenclad®	Merck KGaA	Nemocnice Jihlava, p.o.	No
A 2-year Prospective Study to Assess Health-related Quality of Life in Subjects with Highly-Active Relapsing Multiple Sclerosis Treated with Mavenclad®	Merck KGaA	Nemocnice Pardubického kraje, a.s., Pardubická nemocnice	No
A Multinational, Multicenter, Randomized, Phase 3 Study of Tesetaxel plus a Reduced Dose of Capecitabine versus Capecitabine Alone in Patients with HER2 Negative, Hormone Receptor Positive, Locally Advanced or Metastatic Breast Cancer Previously Treated with a Taxane	Odonate Therapeutics, Inc.	Multiscan s.r.o.	No
A double-blind, randomized, placebo-controlled, multiple-dose, multi-centre safety and efficacy study of co-administration of tesofensine/metoprolol in subjects with Prader-Willi syndrome (PWS) "Second 12 weeks open label extension"	Saniona A/S	Fakultní nemocnice v Motole	No
A Randomized, Double-Blind, Placebo-Controlled, Dose Finding Study to Evaluate the Efficacy and Safety of Padsevonil as Adjunctive Treatment of Focal-Onset Seizures in Adult Subjects with Drug-Resistant Epilepsy	UCB Biopharma SPRL	FORBELI s.r.o.	No
A Randomized, Double-Blind, Placebo-Controlled, Dose Finding Study to Evaluate the Efficacy and Safety of Padsevonil as Adjunctive Treatment of Focal-Onset Seizures in Adult Subjects with Drug-Resistant Epilepsy	UCB Biopharma SPRL	INEP medical s.r.o.	No
A randomized, double-blinded, placebo-Controlled, single and multiple ascending dose study to evaluate the safety, tolerability, pharmacokinetics and pharmacodynamics of BMS-986224 in healthy subjects and chronic heart failure patients with reduced ejection fraction	Bristol-Myers Squibb International Corporation	Všeobecná fakultní nemocnice v Praze	No
A Multicenter, Randomized, Double-Blind, Placebo-Controlled Phase III Study to Evaluate the Efficacy and Safety of Elafibranor in Patients with Non-Alcoholic Steatohepatitis (NASH) and fibrosis.	Genfit SA	Fakultní nemocnice Brno	No
A Phase 3 Randomized, Multicenter Study of Subcutaneous vs. Intravenous Administration of Daratumumab in Subjects With Relapsed or Refractory Multiple Myeloma	Janssen-Cilag International N.V.	Fakultní nemocnice Olomouc	No
Influenza vaccination After Myocardial Infarction (IAMI trial). A multicenter, prospective, randomized controlled clinical trial based on national angiography and angioplasty registries.	Örebro University Hospital	Fakultní nemocnice u sv. Anny v Brně	No
Influenza vaccination After Myocardial Infarction (IAMI trial). A multicenter, prospective, randomized controlled clinical trial based on national angiography and angioplasty registries.	Örebro University Hospital	Fakultní nemocnice Královské Vinohrady	No
A Phase III, Observer-Blind, Randomized, Non-influenza Vaccine Comparator-Controlled, Parallel-Group, Multi-Country Study in Children Aged 6-35 Months to Assess the Safety and Efficacy of Abbott's Candidate Quadrivalent Influenza Vaccine.	Abbott Biologicals B.V.	MUDr. Iva Madejová, praktický lékař pro děti a dorost	No

Safety and efficacy of ex-vivo expanded allogeneic γδ T-lymphocytes (OmnImmune®) in patients with active relapsed or refractory acute myeloid leukaemia (AML) who are not eligible for or do not consent with high dose salvage chemotherapy and/or allogeneic Haematopoietic Cell Transplantation (HCT). A dose escalation, open-label, phase I study.	TC BioPharm, Ltd.	Ústav hematologie a krevní transfuze	No
An Open-Label, Single-Arm Study of The Efficacy, Safety, and Pharmacokinetic Behavior of Leuprolide Mesylate Injectable Suspension (LMIS 25 mg) in Subjects with Prostate Cancer	Foresee Pharmaceuticals Co., Ltd.	Uromedical Center s.r.o.	No
A Multicenter, Randomized, Double-Blind, Placebo-Controlled Phase III Study to Evaluate the Efficacy and Safety of Elafibranor in Patients with Non-Alcoholic Steatohepatitis (NASH) and fibrosis.	Genfit SA	Research Site s.r.o.	No
A Multicenter, Randomized, Double-Blind, Placebo-Controlled Phase III Study to Evaluate the Efficacy and Safety of Elafibranor in Patients with Non-Alcoholic Steatohepatitis (NASH) and fibrosis.	Genfit SA	KLIN MED s.r.o.	No
The Efficacy and Safety of Intra-Arterial Administration of REX-001 to Treat Ischaemic Ulcers in Subjects with Critical Limb Ischaemia Rutherford Category 5 and Diabetes Mellitus: A Pivotal, Placebo-Controlled, Double-Blind, Parallel-Group, Adaptive Trial	Rexgenero Limited	Fakultní nemocnice Ostrava	No
The Efficacy and Safety of Intra-Arterial Administration of REX-001 to Treat Ischaemic Rest Pain in Subjects with Critical Limb Ischaemia Rutherford Category 4 and Diabetes Mellitus: A Pivotal, Placebo-Controlled, Double-Blind, Parallel-Group, Adaptive Trial	Rexgenero Limited	Fakultní nemocnice Ostrava	No
A PROSPECTIVE, RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED PHASE 3 STUDY OF VGX-3100 DELIVERED INTRAMUSCULARLY FOLLOWED BY ELECTROPORATION WITH CELLECTRA™ 5PSP FOR THE TREATMENT OF HPV-16 AND/OR HPV-18 RELATED HIGH GRADE SQUAMOUS INTRAEPITHELIAL LESION (HSIL) OF THE CERVIX	Inovio Pharmaceuticals, Inc.	Všeobecná fakultní nemocnice v Praze	No
A Multinational, Multicenter, Randomized, Phase 3 Study of Tesetaxel plus a Reduced Dose of Capecitabine versus Capecitabine Alone in Patients with HER2 Negative, Hormone Receptor Positive, Locally Advanced or Metastatic Breast Cancer Previously Treated with a Taxane	Odonate Therapeutics, Inc.	Všeobecná fakultní nemocnice v Praze	No
A Placebo-Controlled, Double-Blind, Parallel-Group, 24-Month Study to Evaluate the Efficacy and Safety of E2609 in Subjects with Early Alzheimer's Disease	Eisai Ltd.	AD71 s.r.o.	No
A Placebo-Controlled, Double-Blind, Parallel-Group, 24-Month Study to Evaluate the Efficacy and Safety of E2609 in Subjects with Early Alzheimer's Disease	Eisai Ltd.	BRAIN-SOULTHERAPY s.r.o.	No
Placebo-controlled efficacy and safety trial of intravenous neridronic acid in subjects with complex regional pain syndrome (CRPS)	Grünenthal GmbH	CCR Ostrava, s.r.o.	No
A Phase 3, multicenter, randomized, double-blind, placebo-controlled, parallel-design study to assess the efficacy, safety, and tolerability of AVP-786 (deudextromethorphan hydrobromide [d6-DM]/quinidine sulfate [Q]) for the treatment of agitation in patients with dementia of the Alzheimer's type	Avanir Pharmaceuticals Inc.	Neuropsychiatrie s.r.o.	No
A Phase 3, Randomized, Double-blind, Controlled Study Evaluating the Efficacy and Safety of VX-445 Combination Therapy in Subjects With Cystic Fibrosis Who Are Heterozygous for the F508del Mutation and a Minimal Function Mutation (F/MF)	Vertex Pharmaceuticals Incorporated	Fakultní nemocnice v Motole	No
MONITORING STUDY OF MODERATE TO SEVERE ULCERATIVE COLITIS PATIENTS PREVIOUSLY ENROLLED IN ETROLIZUMAB PHASE II/III STUDIES	F. Hoffmann-La Roche Ltd	ISCARE I.V.F. a.s.	No
Inspection of ethics commitee	-	Ethics commite, Ústav hematologie a krevní transfuze	No
Prevention and treatment of relapse in myeloid and lymphoid neoplasia after allogeneic transplantation with NK cell	Fakultní nemocnice Plzeň	Fakultní nemocnice Plzeň - investigator site inspection	No