Clinical Trials Registry at EDA

SN	Submission Date	Study Code (Specified as per the submitted protocol)	Sponsor/ CRO	Study Title	Study Phase (I, II, III, or IV)	Sites Activation Date "At which the clinical trials will be conducted in Egypt"	Status/Date: - Approved - Recruiting - Recruitment completion - Completed - Withdrawn - Suspended - Terminated	Conditions / Therapeutic Area	Interventions Used IMPs & its type (Biological, Pharmaceutical, Innovative, Herbal, or medical device)
1-	27\12\2018	M15-991	Sponsor: Abbvie	A multi-center, randomized, double-blind, placebo-controlled induction study to assess the efficacy and safety o Risankizumab in subjects with moderately to severely active Crohn's disease who failed prior biologic treatment	III	1- CRC, faculty of medicine, Alexandria university 2- CRC, faculty of medicine, Alexandria university 3- Faculty of medicine, Cairo university 4- MASRI-CRC, Ain	Approved 26/03/2019 Completed 03/11/2021	moderately to severely active Crohn's disease who failed prior biologic treatment	(Biological) Risankizumab

Color Indicator	
Green	Biological
Blue	Pharmaceutical
Orange	Medical Device
Gray	Innovative
Red	Herbal

Registry updated 13/01/2026

						Shams University 5- National hepatology and tropical medicine institute 6- Faculty of medicine, Zagazig university			
2-	27\12\2018	M16-000	Sponsor: Abbvie	A Multicenter, Randomized, Double Blind, Placebo-Controlled 52-Week Maintenance and an Open-Label Extension Study of the Efficacy and Safety of Risankizumab in Subjects with Crohn' Disease who respond to induction treatmen in M16-006 or M15-	III	1- Two sites at Faculty of Medicine, CRC, Alexandria University	Approved 26/03/2019 Recruitment completion	Crohn's disease	(Biological) Risankizumab

				991; or completed M15-989					
3-	28\02\2019	M16-066	Sponsor: Abbvie	A Multicenter, Randomized, Double Blind, Placebo-Controlled 52-Week Maintenance and an Open-Label Extension Study of the Efficacy and Safety of Risankizumab in Subjects with Ulcerative Colitis	III	1- Faculty of medicine, CRC, Alexandria University 2- CRC, Alexandria University 3- Air Force Specialized Hospital Research 4- National Liver Institute, Menoufia University	Approved 10/06/2019 Recruitment completion	Ulcerative Colitis	(Biological) Risankizumab
4-	28\02\2019	M16-067	Sponsor: Abbvie	Multicenter randomized double-blind placebo-controlled induction study to evaluate the efficacy and safety o Risankizumab in	III	1- CRC, faculty of medicine, Alexandria University 2- National Liver Institute,	Approved 10/06/2019 Completed: 30/11/2023	Active ulcerative colitis.	(Biological) Risankizumab

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				subjects with moderately to severely active ulcerative colitis.		Menoufia University 3- Air Force Specialized Hospital 4- Faculty of Medicine, CRC, Alexandria University			
5-	07/05/2019	QGE031	Sponsor: Novartis	A Multicenter, Randomized, double-blind active and placebo-controlled study to investigate the efficacy and safety of Ligelizumab in the treatment of chronic spontaneous urticaria in adolescents and adults in adequately	III	1- Faculty of medicine, Alexandria university 2- Faculty of medicine, Ain Shams University	Withdrawn 31/08/2020	Chronic spontaneous Urticaria	(Biological) Ligelizumab

				controlled with H1 antihistamines					
6-	18/09/2019	ARTEMIS-DM "LPS15396"	Sponsor: SANOVI	A multicenter, multinational, prospective, interventional, single arm, Phase IV study evaluating the clinica efficacy and safety o 26 weeks of treatmer with insulin glargine 300 U/mL (Gla-300) in patients with Type 2 diabetes mellitus uncontrolled on basa insulin	IV	1- Faculty of medicine, Alexandria university 2- CRC, Alexandria university 3- GOTHI 4- Faculty of medicine, Menoufia university 5- Faculty of medicine, Ain Shams university	Approved 09/02/2020 Withdrawn	Type 2 diabetes mellitus	(Biological) Insulin glargine "Toujeo"
7-	18/11/2019	STAND	Sponsor: Novartis	A phase II, multicenter, randomized, open label, two arm study comparing the effect of crizanlizumab+ SOC alone on renal	II	1- Abu El Resh Children Hospital	Approved 05/05/2020 Withdrawn 03/08/2021	Sickle cell anemia	(Biological) Crizanlizumab

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				function in sickle cel disease patients ≥16 years with chronic kidney disease due to sickle cell nephropathy					
8-	24/03/2020	STEAD FAST	Sponsor: Novartis	A Phase III, multicenter, double-blind study to assess efficacy and safety o two doses of crizanlizumab vs placebo with or without hydroxyurea / hydroxycarbamide therapy, in adolescen and adult sickle cell disease patients with vaso-occlusive crisis	III	1- Faculty of medicine, Alexandria university 2- Faculty of medicine, Ain Shams university	Approved 20/02/2020 Withdrawn 03/08/2021	Sickle cell anemia	(Biological) Crizanlizumab
9-	30/03/2020	WA40404	Sponsor: Roche	A Phase III b Multicenter, Randomized, double-blind, Placebo-controlled study to evaluate the efficacy	IIIb	1- Sayed Galal Hospital 2- Faculty of medicine, Alexandria university	Approved 23/08/2020 Withdrawn 25/08/2021	Primary progressive multiple sclerosis	(Biological) Ocrelizumab

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				and safety of Ocrelizumab in adult with primary progressive Multiple Sclerosis		3- CRC, MASRI, Ain Shams University			
10-	14\09\2020	1368-0025	Sponsor: Boehringer Ingelheim	Open label long term extension study to assess the safety and efficacy of BI655130 treatment in patients with generalized pustular psoriasis	IIb	1- Dermatology department, faculty of medicine, Alexandria university hospital	Approved 18/05/2021 Withdrawn 31/10/2021	Generalized pustular psoriasis	(Biological) Spesolimab
11-	21/09/2020	05-Gam-COVID-Vac-2020	Sponsor: Russian Direct Investment Fund (RDIF)	A Phase III, randomized, double blind, placebo-controlled trial to evaluate immunogenicity and safety of the Gam-COVID-Vac combined vector vaccine in prophylactic treatment for SARS-	III	1- National liver institute, Menoufia university 2- CRC, faculty of medicine, Alexandria university 3- CRC, MASRI, Ain Shams University	Withdrawn 12/06/2022	COVID-19 prophylaxis	(Biological) Russian Gam-COVID-Vac Combine vector vaccine

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				COV-2 infection in Egypt					
12-	22/09/2020	CNBG2020003SQ	Sponsor: China National Biotec Group company limited Wuhan institute of biological products Co. Ltd Beijin institute of biological products Co.Ltd	Multicenter, Randomized, Double blind, parallel placebo controlled, Phase III clinical tria to evaluate the protective efficacy, safety and immunogenicity of Inactivated SARS-COV-2 Vaccines in healthy population aged 18 years old and above	III	1- Vacsera Health care facility 2- Ktameya medical center	Approved 28/03/2022 Completed 31/07/2022	COVID-19 Prophylaxis	(Biological) Inactivated SARS-COV-1 Vaccine
13-	13/04/2021	D910DC00001 (Emerald-2)	Sponsor: AstraZeneca CRO: IQVIA	A phase 3 randomized double blind placebo controlled multicentr study of durvalumab monotherapy or in	III	1- CRC, Faculty of medicine, Alexandria University hospital	Approved 12/12/2021 Recruitment completion	Hepatocellular carcinoma patients at high risk of recurrence after curative	(Biological) Durvalumab\ Bevacizumab

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				combination with bevacizumab as adjuvant therapy in patients with hepatocellular carcinoma who are at high risk of recurrence after curative hepatic resection or ablation		2- National Liver Institute-Menoufia University 3- National Hepatology & Tropical Medicine Research Institute 4- Air Force specialized Hospital 5- Faculty of medicine, Assuit University		hepatic resection or ablation	
14-	19/05/2021	01-Sputnik-Light-2021	Sponsor: Human vaccine LLC (Global), Russian ministry	A phase III, randomized, double-blind, placebo-controlled international multi-site clinical trial in parallel assignment t	III	1- National hepatology and tropical medicine center 2- Katemeya medical center	Approved 24/08/2021 Completion of study visit 31/08/2022	COVID-19 Prophylaxis	(Biological) Sputnik Light vector vaccine

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			of healthcare – Gamalya (Local) CRO: PDC	evaluate efficacy, immunogenicity and safety of the Sputnik Light vector vaccine in adults in the SARS-Cov-2 infection prophylacti treatment					
15-	25/05/2021	KATE-3	Sponsor: Roche	A randomized, multi-center, double blind, placebo-controlled phase III study of the efficacy and safety o Trastuzumab Emtansine in combination with Atezolizumab or placebo in Pts with HER2-positive and PD-L1- positive locally advanced or metastatic breast cancer who have received prior	III	1- Faculty of medicine, Kasr Al-Ainy hospital 2- Shefaa Al-Orman hospital 3- Baheya Hospital	Approved 05/12/2021 Withdrawn 19/12/2022	HER2-positive and PD-L1-positive locally advanced or metastatic breast cancer	(Biological) Trastuzumab Emtansine/ Atezolizumab

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				Trastuzumab + Atezolizumab and Taxane- based therapy					
16-	27/05/2021	CAIN457 P12301	Sponsor: Novartis	A randomized, double blind, placebo-controlled, parallel group, phase III multi-center study of intravenous Secukinumab to compare efficacy at 16 weeks with placebo and to assess safety and tolerability up to 52 weeks in subjects with active ankylosis spondylitis of non-radiographic axial spondylo arthritis	III	1-CRC, Faculty of medicine, Alexandrian university	Withdrawn 03/11/2021	Active ankylosing spondylitis	(Biological) Secukinumab
17-	05/08/2021	TG2101V 01	Sponsor: Livzon mabpharm Inc.	A Global, Multi-Center, Randomized, Double-Blind, Placebo-Controlled,	III	1-National Hepatology and Tropical	Withdrawn 16/01/2022	COVID-19 Prophylaxis	(Biological) Recombinant SARS-CoV-2

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				Phase III Clinical Study to Evaluate the Efficacy, Safety and Immunogenicity of Recombinant SARS-CoV-2 Fusion Protei Vaccine (V-01) in Adults Aged 18 Years and Older",		Medicine Research Institute (NHTMRI)			Fusion Protein Vaccine (V-01)
18-	18/08/2021	MO42541	Sponsor: Roche	A phase III, open label, randomized study of Atezolizumab with Lenvatinib or Sorafenib versus Lenvatinib or sorafenib alone in hepatocellular carcinoma previously treated with Atezolizumab and Bevacizumab	III	1- Air force specialized hospital	Approved 02/02/2022 Recruitment completion	Hepatocellular carcinoma	(Biological) Atezolizumab/ Lenvatinib/ Sorafenib
19-	02/09/2021	COVID_ VACC_1	Sponsor: National	A Phase 1 Clinical Trial to Evaluate the Safety,	I	1- National research center	Approved 09/11/2021	Covid-19 Prophylaxis	(Biological)

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			research center CRO: Clinmax	Tolerability, and Immunogenicity of Inactivated SARS-CoV-2 Vaccine Against COVID-19 in Healthy Adults			Suspended 09/12/2021		Inactivated SARS-CoV-2 Vaccine
20-	17/01/2022	SPHINX-EGYPT SPHINX2 2122020	Sponsor: - EVA PHARMA - VSVRI - supreme council of university hospitals - Ministry of higher education and scientific research CRO: Dataclin	Safety and Immunogenicity Study of EgyVax Vaccine Candidate for Prophylaxis of SARS-CoV-2 Infection (COVID-19)	I	1- Al-Manial specialized university Hospital, Cairo university hospitals	Approved 03/02/2022 Database lock 26/09/2023	Covid-19 Prophylaxis	(Biological) EgyVax
21-	04/11/2021	GBT2104-131	Sponsor:	A randomized double blinded	III	1- Faculty of medicine,	Approved 14/06/2022	sickle cell disease	(Biological)

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			Global blood therapeutics Inc. \ Pfizer CRO: MCT	placebo controlled multicentre study to access the safety and efficacy of Inclacumab in participants with sickle cell disease experiencing Vaso-occlusive crisis		Mansoura University 2- Faculty of medicine, Zagazig University 3- MASRI-CRC, Faculty of medicine, Ain Shams University hospital 4- CRC, Alexandria University 5- Pediatric hematology department, Alexandria University 6- CRC, faculty of medicine, Cairo University, Abo	Completed: 30/05/2025	patients with Vaso-occlusive crisis	Inclacumab

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						El-Resh Hospital 7- CRC, Cairo University 8- Hematology department, Cairo University hospital			
22-	04/01/2022	GBT2104-132	Sponsor: Global blood therapeutics Inc.\ Pfizer CRO: MCT	A Randomized, Double-blind, Placebo-controlled, Multicenter Study of a Single Dose of Inclacumab to Reduce Re-admission in Participants with Sickle Cell Disease and Recurrent Vaso-occlusive Crises (GBT-132)	III	1- Faculty of medicine, Mansoura University 2-. Faculty of medicine, Zagazig University 3- MASRI, CRC, Ain Shams University 4- Hematology unit, Internal medical department, CRC, faculty of medicine	Approved 14/06/2022 Withdrawn 29/06/2023	Sickle cell disease patients with Vaso-occlusive crisis	(Biological) Inclacumab

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						Alexandria University hospital 5- Hematology department, Alexandria University hospital 6- Cairo University, Abo El-Resh Hospital 7- CRC, Cairo University 8- Cairo University, Hematology department.			
23-	28/11/2021	GBT2104-133	Sponsor: Global blood therapeutics Inc.\ Pfizer	An Open-label Extension Study to Evaluate the Long-term Safety of Inclacumab Administered to Participants with Sickle Cell Disease	III	1- Faculty of medicine, Mansoura University 2- Faculty of medicine, Zagazig University	Approved 14/06/2022 Withdrawn 17/12/2023	sickle cell disease	(Biological) Inclacumab/ Placebo

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			CRO: MCT	Who Have Participated in an Inclacumab Clinical Trial		3- MASRI, CRC, Ain Shams University 4- Hematology unit, Internal medical department, CRC, faculty of medicine Alexandria University hospital 5- Hematology department, Alexandria University hospital 6- Cairo University, Abo El-Resh Hospital 7- CRC, Cairo University 8- Cairo University,			

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						Hematology department.			
24-	08\06\2022	Consonance-MN39159	Sponsor: F. HOFFMANN-LA ROCHE LTD CRO: Roche Egypt LLC & IQVIA (for monitoring activities only)	An open-label, single-arm 4-year study to evaluate effectiveness and safety of ocrelizumab treatment in patients with progressive multiple sclerosis	III	1- CRC, Faculty of Medicine, Alexandria university, CRC 2- MASRI-CRC, faculty of medicine, Ain Shams university hospital	Approved 20/09/2022 Recruitment completion	Progressive multiple sclerosis	(Biological) Ocrelizumab
25-	09\02\2022	20200404 (IMBCAM	Sponsor: Institute of Medical Biology Chinese Academy of	A randomized double-blinded placebo-controlled Phase III clinical trial of SARS-COV-2 vaccine inactivated	III	1- Katameya Medical Center 2- National Hepatology and tropical medicine institute	Withdrawn 24/02/2022	Covid-19 Prophylaxis	(Biological) Inactivated SARS-COV-2 vaccine

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			Medical Sciences CRO: PDC	(Vero cell) in adult aged 18 years and above					
26-	10/05/2022	TRISTARDS-0135-0347	Sponsor: Boehringer Ingelheim CRO: MCT	The TRISTARDS trial -ThRombolys is Therapy for ARDS A Phase IIb/III operationally seamless, open-label, randomized, sequential, parallel-group adaptive study to evaluate the efficacy and safety of daily intravenous alteplase treatment given up to 5 days on top of standard of care (SOC) compared with SOC alone, in	IIb/III	1- National Hepatology and Tropical Medicine Research Institute 2- Abbasia Fever Hospital 3- Imbaba Fever Hospital	Withdrawn 20/07/2022	Respiratory distress syndrome (ARDS) triggered by COVID-19	(Biological) Alteplase

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				patients with acute respiratory distress syndrome (ARDS) triggered by COVID-19.					
27-	14/08/2022	CAIN457 A2310	Sponsor: Novartis CRO: MCT	A randomized, double-blind, placebo- and active controlled multicenter trial to demonstrate efficacy of subcutaneous Secukinumab compared to placebo and etanercept (in a single blinded arm) after twelve weeks of treatment, and to assess the safety, tolerability, and long-term efficacy in subjects from 6 to less than 18	III	1- CRC, Faculty of Medicine, Alexandria university hospital 2- Dermatology department, faculty of Medicine, Ain Shams University hospital	Approved 04/12/2022 Early terminated by sponsor 31/03/2023	Treatments of severe chronic plaque psoriasis	(Biological) Secukinumab

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				years of age with severe chronic plaque psoriasis					
28-	08/11/2022	SCTV01E -MRCT-1	Sponsor: Sinocelltech CRO: PDC	A randomized double blind positive controlled phase III clinical trial to evaluate the efficacy and safety of SCTV01E (a covid-19alpha/beta/delta/ omicron variants s-trimmer vaccine) in population previously unvaccinated with COVID-19 vaccine and aged ≥18	III	1- Katemya Medical Center 2- Egyptian Liver research institute and hospital	Withdrawn 14/01/2023	COVID-19 prophylaxis	(Biological) SCTV 01E (a covid-19 alpha/beta/delta/ omicron variants s-trimmer vaccine) (Biological)
29-	06/06/2023	FUZION CNTO1959CRD	Sponsor: Janssen CRO: MCT	A Phase 3, Randomized, Placebo-controlled, Parallel-group, Multicenter Study to Evaluate the	III	1- National Hepatology Tropical Medicine Research Institute	Approved 13/08/2023 Recruiting	Fistulizing perianal Crohn's disease	Guselkumab (Biological)

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				Efficacy and Safety of Guselkumab in Participants with Fistulizing, Perianal Crohn's Disease "FUZION CD"		2- CRC, faculty of medicine Alexandria university hospital, (two sites) 3- Department of internal medicine, El Kasr Al Aini, Cairo University 4- MASRI CRC, faculty of medicine, Ain Shams University Hospital			
30-	14/05/2023	MP-ADA1-01	Sponsor: Minapharm CRO: CRS Clinical Research	A Phase I, randomized, double-blind, 2-arm, parallel group trial to compare pharmacokinetics of Adessia with EU-authorized	I	1- CRS clinical research services, Berlin GmbH 2- CRS clinical research services,	Approved 10/08/2023 Completed	Inflammatory disease (Biosimilar to Humira)	Adessia (Biological)

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			Services Berlin GmbH	Humira in healthy male and female participants"		Mannheim GmbH			
31-	04/05/2023	MOM-M281-006	Sponsor: Janssen CRO: MCT	Efficacy and Safety of M281 in Adults with Warm Autoimmune Hemolytic Anemia: A Multicenter, Randomized, Double-blind, Placebo-controlled Study with a Long-term Open-label Extension"	II\III	1- National Cancer Institute, Cairo university 2- Oncology center, Mansoura University Hospital 3- Department of internal medicine, Al Kasr al Eini, Cairo university 4- Naser institute hospital for research and treatment 5- CRC, faculty of medicine, Alexandria university Hospital	Approved 19/07/2023 Early Terminated by the sponsor 21/02/2025	Warm Autoimmune Hemolytic Anemia	M281 (Biological)

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						6- CRC, faculty of medicine, Ain shams university Hospital			
32-	09\10/2023 shift to amendment submission 26\12\2023	EMERALD-3) D910VC00001	Sponsor: AstraZeneca CRO: IQVIA	A Phase III, Randomized, Open-Label, Sponsor-Blinded, Multicenter Study of Durvalumab in Combination with Tremelimumab ± Lenvatinib Given Concurrently with Transarterial Chemoembolization (TACE) Compared to TACE Alone in Patients with Locoregional Hepatocellular Carcinoma (EMERALD-3)	III	1- Air Force specialized hospital 2- Oncology department, Faculty of medicine, Alex University 3- Egyptian liver Hospital National Hepatology and Tropical Medicine Research Institute (NHTMRI) 4- Shifa El orman Hospital	Approved 08/02/2024 Recruiting	Locoregional Hepatocellular Carcinoma	(Biological) Durvalumab / Tremelimumab/ Lenvatinib /TACE

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33-	not submitted officially	CERE-CAP	Investigator-initiated	Efficacy of Cerebrolysin as an adjuvant therapy following mechanical thrombectomy in patients with large vessels occlusion stroke	III	1- Neurology and psychiatry department, Ain Shams University Hospital	Terminated (by EDA) (15/01/2024)	occlusion stroke	(Biological) CEREBROLYSIN solution for IM or IV injection/ concentrate for solution for I.V. infusion
34-	14/12/2023	BCD-178	Sponsor: JSC BIOCAD CRO: Dataclin	A Double-Blind, Randomized Clinical Study of the Efficacy and Safety of BCD-178 and Perjeta® as Neoadjuvant Therapy of HER2-Positive Breast Cancer	III	1- Faculty of Medicine, Alexandria University 2- Faculty of Medicine, Cairo University	Approved: 22/04/2024 Withdrawn: 25\12\2024	Her-2 positive breast cancer	Biological BCD-178
35-	08/01/2024	SerpinPc 102	Sponsor: Apcintex CRO: MCT	A Global, Open-label, Adaptive Design Study to Investigate the Efficacy and Safety of SerpinPC in	IIb	1- Ain Shams University Medical Research Institute (MASRI)	Conditional Approved 13/06/2024 Final Approval 31/10/2024	Hemophilia A or Moderately Severe to Severe	Biological SerpinPC 102

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				Subjects with Severe Hemophilia A or Moderately Severe to Severe Hemophilia B (AP-0102)			Withdrawn: 16/01/2025	Hemophilia B	
36-	08/01/2024	SerpinPC103	Sponsor: Apcintex CRO: MCT	A Global, Open-label Study to Investigate the Efficacy and Safety of SerpinPC in Subjects with Hemophilia B with Inhibitors (AP-0103)	IIb	1- Ain Shams University Medical Research Institute (MASRI)	Conditional Approved 13/06/2024 Final Approval 31/10/2024 Withdrawn: 16/01/2025	Hemophilia B with Inhibitors	Biological Serpin PC 103
37-	08/02/2024	D9185C00001” TILIA’	Sponsor: AstraZenca CRO: IQVIA	A Phase III, Multicenter, Randomized, Double-bind, Parallel-group, Placebo-Controlled study to evaluate the efficacy and safety of	III	1-Air Force specialized Hospital 2-Ain Shams University Medical Research Institute (MASRI-CRC)	Approved: 04/08/2024 Withdrawn: 30/01/2025	Patients hospitalized for viral lung infection	Biological Tozoralimab

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				Tozoralimab (MEDI3506) in patients hospital;ized for viral lung infection requiring supplemental oxygen		3-CRC, Alexandria University Hospital			
38-	16/01/2025	GA45329-Ametrine 1	Sponsor: Roche	A Phase III, Multicenter, Double-Blind, Placebo-Controlled, Treat-Through Study to Assess the Efficacy and Safety of Induction and Maintenance Therapy With RO7790121 In Patients with Moderately to Severely Active Ulcerative Colitis	III	1- Clinical Research Center, Internal Medicine Faculty of Medicine, Alexandria University	Approved: 10/03/2025 Withdrawn: 10/09/2025	Patients with Moderately to Severely Active Ulcerative Colitis	Biological RO7790121

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39-	16/01/2025	GA45330-Ametrine 2	Sponsor: Roche	A Phase III, Multicenter, Double-Blind, Placebo-Controlled Study to Assess the Efficacy and Safety of Induction Therapy with RO7790121 In Patients with Moderately to Severely Active Ulcerative Colitis	III	1- Clinical Research Center, Internal Medicine Faculty of Medicine, Alexandria University	Approved: 10/03/2025 Withdrawn: 10/09/2025	Patients with Moderately to Severely Active Ulcerative Colitis	Biological RO7790121
40-	11\05\2025	GA45331	Sponsor: Roche	A Phase III Multicenter Double Blind Placebo Controlled Treat through study to assess the efficacy and safety of Induction and Maintenance therapy with RO7790121 in patients with	III	1-Clinical Research Center, Faculty of Medicine, Alexandria University Hospital 2-Air Force Specialized Hospital 3-National Liver Institute,	Approved: 01\06\2025	Crohn's Disease	Biological RO7790121

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				moderately to severely active Crohn's disease		Menoufia University			
41-	20/05/2025	M20-465	Sponsor: Abbvie	A Phase III Multicenter, Randomized, Double-Blind Placebo-Controlled Study to Evaluate the Efficacy and Safety of Lutikizumab in Adult and Adolescent Subjects with Moderate to Severe Hidradenitis Suppurativa	III	1.Air Force Specialized Hospital 2. CRC, Faculty of Medicine, Alexandria University (2 sites) 3. MASR CRC, Faculty of Medicine, Ain Shams University	Approved: 08/10/2025	Moderate to Severe Hidradenitis Suppurativa	Biological Lutikizumab
42-	18/09/2025	GA45332	Sponsor: Roche	A Phase III, Multicenter, Double-Blind< Placebo-Controlled Study to assess the efficacy and safety of induction therapy with RO7790121	III	1-Clinical Research Center, Faculty of Medicine, Alexandria University Hospital	Approved: 20/10/2025	Crohn's Disease	Biological RO7790121

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				in patients with moderately to severely active Crohn's disease		2-Air Force Specialized Hospital 3-National Liver Institute, Menoufya University			
43-	17/12/2020	CEGA230 B2404	Sponsor: Novartis CRO: MCT	A Phase IV Multicenter Open Label Study to Determine the Safety, Tolerability and Clinical Outcomes Following Oral Administration of Egaten (Triclabandazole) in Patients 6 Years of Age or Older with Fascioliasis (Egaten)	IV	1-Cairo University, Al Mounira Children Hospital, Pediatric Hepatology Unit. 2-Alexandria University, Faculty of Medicine, Clinica Research Center.	Approved 12/04/2021 Recruitment Completion	Fascioliasis	(Pharmaceutical) Triclabandazole (Egaten)
44-	22/12/2020	CLEE011 A3201C	Sponsor: Novartis	A Phase II Randomized Study	II	1-Ain Shams University, Faculty of	Approved 14/10/2021	HER-2 Negative	(Pharmaceutical)

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		RIGHT Choice	CRO: MCT	of the Combination of Ribociclib Plus Goserelin Acetate with Hormonal Therapy Versus Physician Choice Chemotherapy in Premenopausal or Perimenopausal Patients with Hormone Receptor-Positive/HER2-Negative Inoperable Locally Advanced or Metastatic Breast Cancer - RIGHT Choice Study (RIGHT Choice)		Medicine, Clinical Research Center, (MASRI – CRC) 2-Baheya Hospital Research Center 3-Cairo University, NEMROCK 4-Nasser Institute Cancer Center	Completed 08/01/2023	Breast Cancer	Ribociclib Plus Goserelin / Physician Choice Chemotherapy
45-	24/10/2021	M14-430	Sponsor: Abbvie	A Multicenter, Randomized, Double-Blind,	III	1-Air Force Specialized Hospital	Approved 07/07/2022	Chron's Disease	(Pharmaceutical)

Color Indicator	
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Blue	Pharmaceutical
Orange	Medical Device
Gray	Innovative
Red	Herbal

				Placebo-Controlled Maintenance and Long-Term Extension Study of the Efficacy and Safety of Upadacitinib (ABT-494) in Subjects with Crohn's Disease Who Completed the Studies M14-431 or M14-433		2-National Liver Institute Menoufiya University 3-Alexandria University, Faculty of Medicine, Clinical Research Center. 4-Ain Shams University, Faculty of Medicine, Clinical Research Center (MASRI-CRC).	Recruitment Completion		Upadacitinib/ matching placebo
46-	26/10/2021	BO40336 ALINA	Sponsor: Roche	A Phase III, Open-Label, Randomized Study to Evaluate the Efficacy and Safety of Adjuvant Alectinib Versus Adjuvant Platinum-Based Chemotherapy in Patients with	III	1- Cairo University, Kasr Al Eini, Center of Radiation Oncology and Nuclear Medicine.	Approved 16/03/2022 Recruitment Completion	Lung Cancer	(Pharmaceutical) Alectinib / Platinum based Chemotherapy

Color Indicator	
Green	Biological
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Gray	Innovative
Red	Herbal

				Completely Resected Stage Ib (Tumors ≥ 4 Cm) To Stage IIIa Anaplastic Lymphoma Kinase-Positive Non-Small-Cell Lung Cancer (ALINA)					
47-	12/12/2021	Cl_Tr_17 122019 MIRACL E-ALA	Sponsor: EVA Pharma CRO: MARC	A Multicenter, Interventional, Two-Arm, Parallel-Group, Randomized, Double-Blinded, Placebo-Controlled, Phase IV Trial to Evaluate the Efficacy of Alpha-Lipoic Acid in the Treatment of Patients with Symptomatic Diabetic	IV	1- Alexandria University Hospital, Diabetes, Metabolism, and Lipidology Unit, Department of Internal Medicine. 2- Ain Shams University Hospital	Approved 12/10/2022 Completed 11/12/2024	Treatment of Symptomatic Diabetic Polyneuropathy	(Pharmaceutical) Alpha-Lipoic Acid (Thiotacid)/ matching placebo

Color Indicator	
Green	Biological
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Gray	Innovative
Red	Herbal

				Polyneuropathy in Egypt (MIRACLE-ALA)		3- Menoufiya University Hospital 4- Mansoura University, Intrinsic Specialized Hospital. 5- Beni-Suef University Hospital, Diabetes and Endocrinology Unit.			
48-	12/12/2021	MK4482-013 MOVe-Ahead	Sponsor: MSD	A Phase 3 Multicenter, Randomized, Double Blind, Placebo Controlled Study to Evaluate the Efficacy and Safety of MK-4482 for the Prevention of COVID-19 (Laboratory	III	1-Ain Shams University Clinical Research Center (MASRI-CRC). 2-Air Force Specialized Hospital. 3-National Hepatology and Tropical	Approved 18/01/2022 Completed 16/11/2022	Prophylaxis of COVID-19	(Pharmaceutical) Molnupiravir/ matching placebo

Color Indicator	
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Gray	Innovative
Red	Herbal

				Confirmed SARS-COV 2 Infection with Symptoms) in Adults. (MOVe-Ahead)		Medicine Research Institute. 4-Imbaba Fever Hospital. 5-National Center for Allergies and Chest Imbaba			
49-	30/03/2022	GBT440-032	Sponsor: GBT (Subsidiary of Pfizer) CRO: CTI	A Phase 3, Randomized, Double-Blind, Placebo-Controlled Study of Voxelotor (GBT440) in Pediatric Participants with Sickle Cell Disease (HOPE Kids 2)	III	1-Ain Shams University Clinical Research Center (MASRI-CRC). 2-Alexandria University Clinical Research Center. 3- Al Mounira Children Hospital, Cairo University, 4-Zagazig University	Approved 31/07/2022 IMP Dosing Pause 02/05/2024 Early Termination by the sponsor 29/09/2024	Sickle Cell Disease	(Pharmaceutical) Voxelotor/ matching placebo

Color Indicator	
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Red	Herbal

						Hospital, Department of Pediatrics.			
50-	18/04/2022	GBT440-034	Sponsor: GBT (Subsidiary of Pfizer) CRO: IQVIA	An Open Label Extension Study of GBT440 Administered Orally to Patients with Sickle Cell Disease who Have Participated in GBT440 Clinical Trials	III	1-Cairo University, Abu El Rich Hospital. 2-Ain Shams University Clinical Research Center (MASRI-CRC) 3-Alexandria University Clinical Research Center 4-Zagazig University Hospital, Department of Pediatrics.	Approved 02/08/2022 Early Termination by the sponsor 30/09/2024	Sickle Cell Disease	(Pharmaceutical) Voxelotor
51-	17/05/2022	F901318/0032	Sponsor: F2G	Open Label Single Arm Phase IIb Study of F901318 as Treatment of	IIb	1-Mansoura University Oncology center	Terminated (By Sponsor) 24/07/2022	Invasive Fungal Infection	(Pharmaceutical) Olorofim

Color Indicator	
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			CRO: IQVIA	Invasive Fungal Infections Due to Lomentospora Prolificans, Seedosporium Spp., Aspengillus Spp., & other Resistant Fungi in Patients Lacking Suitable Alternative		2-Alexandria University, Clinical Research Center 3-Nasser Institute 4-Ain Shams University Clinical Research Center, (MASRI – CRC) 5-Air Force specialized Hospital 6-National Cancer Institute 7-Cairo University Kasr Al-Eini, Hospital			
52-	12/06/2022	CLSYN.1702 (OASIS-9)	Sponsor: Hamilton Health Science	A 2x2 Factorial Randomized Controlled Trial of CoLchicine and	III/IV	1-Mansoura University Hospital	Approved 24/07/2022 Completed	STEMI/Non-STEMI Myocardial Infarction	(Pharmaceutical) Colchicine, Spironolactone/

Color Indicator	
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			CRO: Clinmax	spironolactonE in Patients With myocARdial infarction/SYNERGY Stent Registry – Organization to Assess Strategies for Ischemic Syndromes 9 (OASIS-9)		2-Suez Canal University Hospital 3-Fayoum General Hospital 4-Tamia Central Hospital 5-El Kharga Specialized Hospital 6-National Heart Institute	01/08/2024		matching placebo
53-	15/06/2022	20140106	Sponsor: Onyx Pharmaceuticals (Subsidiary of Amgen) CRO: IQVIA	Phase 1b/2 Study of Carfilzomib in Combination with Induction Chemotherapy in Children with Relapsed or Refractory Acute Lymphoblastic Leukemia	Ib/II	1-Children's Cancer Hospital 57357	Approved 23/08/2022 Withdrawn 19/06/2023	Relapsed or Refractory Acute Lymphoplastic Leukemia	(Pharmaceutical) Carfilzomib
54-	18/07/2022	AG348-C-020	Sponsor: Agios	A Phase 2/3, Double-Blind, Randomized,	II/III	1-Alexandria University	Approved 27/09/2022	Sickle Cell Disease	(Pharmaceutical) Mitapivat /

Color Indicator	
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			CRO: MCT	Placebo-Controlled, Multicenter Study to Evaluate the Efficacy and Safety of Mitapivat in Subjects with Sickle Cell Disease		Clinical Research Center 2-Zagazig University Hospital 3-Cairo University Hospital 4-Mansoura University Hospital 5-Ain Shams University Clinical Research Center (MASRI-CRC)	Withdrawn 21/08/2023		matching placebo
55-	26/07/2022	F901318/0041	Sponsor: F2G CRO: IQVIA	A Phase III, Adjudicator-Blinded, Randomised Study to Evaluate the Efficacy and Safety of Treatment with Olorofim Versus Treatment with	III	1-Mansoura University Oncology Center 2-Alexandria University Clinical Research Center	Approved 11/10/2022	Invasive Fungal Disease caused by Aspergillus species	(Pharmaceutical) Olorofim / Ambisome

Color Indicator	
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				Ambisome® Followed by Standard of Care (SOC) in Patients with Invasive Fungal Disease (IFD) Caused by Aspergillus Species		3-Air Force specialized Hospital 4-Ain Shams University, Clinical Research Center (MASRI-CRC) 5-Zagazig University Hospital 6-National Cancer Institute 7-Cairo University Kasr Al Eini Hospital 8-Nasser Institute for Research and Treatment			
56-	27/07/2022	APD334-202	Sponsor: Arena Pharmaceuticals (Subsidiar	A Multicenter Randomized Double Blinded Parallel Group Study to Assess the	III	1-Alexandria University Clinical Research Center	Approved 23/08/2022 Recruitment Completion	Moderately to Severe Active Crohn's Disease	(Pharmaceutical) Etrasimod / matching placebo

Color Indicator	
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			y of Pfizer)	Efficacy and Safety of Oral Etrasimod as Induction and Maintenance Therapy for Moderately to Severe Active Crohn's Disease (Etrasimod)		2-Air Force Specialized Hospital 3-National Liver Institute 4-National Hepatology and Tropical Medicine Research Institute (NHTMRI) 5-Cairo University Kasr Al-Eini Hospital 6-Egyptian Liver Research Institute and Hospital 7-Ain Shams University Hospital 8-Theodor Bilharz	Early Terminated by the sponsor 20/03/2025		

Color Indicator	
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						Research Institute			
57-	07/08/2022	EFC17215 LEAP-2-MONO	Sponsor: Sanofi	A Phase 3, Multicenter, Multinational Randomized Double-Blind Double-Dummy, Active Comparator Study to Evaluate the Efficacy and Safety of Venglustat in Adult and Pediatric Patients with Gaucher Disease Type 3 (GD3) who Have Reached Therapeutic Goals with Enzyme Replacement Therapy (LEAP-2-MONO)	III	1-Alexandria University Hospital Clinical Research Center	Approved 24/10/2022	Gaucher Disease Type 3 (GD3)	(Pharmaceutical) Venglustat/ Cerezyme

Color Indicator	
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58-	15/08/2022	AG348-C-017	Sponsor: Agios CRO: MCT	A Phase 3, Double-blind, Randomized, Placebo-Controlled, Multicenter Study Evaluating the Efficacy and Safety of Mitapivat in Subjects with Non–Transfusion-Dependent Alpha- or Beta-Thalassemia (ENERGIZE)	III	1-Cairo University Hospital 2-Ain Shams University Clinical Research Center MASRI-CRC	Approved 02/11/2022 Withdrawn 26/06/2023	Non-Transfusion-Dependent Alpha or Beta Thalassemia	(Pharmaceutical) Mitapivat / matching placebo
59-	15/08/2022	AG348-C-018	Sponsor: Agios CRO: MCT	A Phase 3, Double-blind, Randomized, Placebo-Controlled, Multicenter Study Evaluating the Efficacy and Safety of Mitapivat in Subjects with Transfusion-Dependent Alpha-	III	1-Cairo University Hospital 2-Ain Shams University Clinical Research Center MASRI-CRC	Approved 02/11/2022 Withdrawn 26/06/2023	Transfusion-Dependent Alpha or Beta Thalassemia	(Pharmaceutical) Mitapivat / matching placebo

Color Indicator	
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				or Beta-Thalassemia (ENERGIZE-T)					
60-	29/08/2022	4202-HEM-301	Sponsor: Forma Therapeutics CRO: MCT	An Adaptive, Randomized, Placebo-Controlled, Double-blind, Multi-center Study of Oral Etavopivat, a Pyruvate Kinase Activator in Patients with Sickle Cell Disease (Hibiscus Study)	III	1- Alexandria University Clinical Research Center 2-Zagazig University Hospital 3-Cairo University Hospital 4-Ain Shams University Clinical Research Center (MASRI-CRC)	Approved 11/12/2022 Recruitment Completion	Sickle Cell Disease	(Pharmaceutical) Etavopivat / matching placebo
61-	29/09/2022	GO42784 LIDERA	Sponsor: Roche CRO: MCT	A Phase III, Randomized, Open-Label, Multicenter Study Evaluating the Efficacy and Safety of Adjuvant	III	1-Alexandria University Hospital 2-Medical Research Institute,	Approved 04/12/2022 Recruitment Completion	Estrogen Receptor−Positive, Her2-Negative Early Breast Cancer	(Pharmaceutical) Giredestrant / Physician Choice of Adjuvant Endocrine Monotherapy

Color Indicator	
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Red	Herbal

				Giredestrant Compared with Physician's Choice of Adjuvant Endocrine Monotherapy in Patients with Estrogen Receptor−Positive, Her2-Negative Early Breast Cancer (LIDERA)		Alexandria University 3-Mansoura University Hospital 4-Cairo University Kasr Al- Ainy Hospital 5-Ain Shams University Demerdash Hospital 6- Dar El Salam Cancer Hospital 7- Sohag Oncology Center			
62-	16/11/2022	(ACTIV-2D/A5407)	Sponsor: Shionogi CRO: IQVIA	A Phase 3, Multicenter, Randomized, Double-Blind, 24-Week Study of the Clinical and Antiviral Effect of	III	1-National Hepatology and Tropical Medicine Research Institute	Approved 31/01/2023 Withdrawn 26/09/2023	Covid-19 treatment	(Pharmaceutical) S-217622 / matching placebo

Color Indicator	
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				S-217622 Compared with Placebo in Non-Hospitalized Participants with COVID-19 (ACTIVE)		2-Ain Shams University Clinical Research Center (MASRI-CRC) 3-Alexandria University Clinical Research Center, 4-Air Force Specialized Hospital 5-National Institute for Chest Allergy and Diseases 6-Imbaba Fever Hospital			
63-	28/11/2022	RBSC2161	Sponsor: Salix pharmaceuticals CRO:	A Phase 2a Randomized, Double-Blind, Placebo-Controlled Study to Characterize the	IIa	1-Cairo University Abu El Rich Hospital. 2-Ain Shams University	Approved 05/02/2023 Withdrawn 06/11/2023	Sickle Cell Disease	(Pharmaceutical) Rifaximin / matching placebo

Color Indicator	
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			IQVIA	Pharmacokinetics and Pharmacodynamics of Rifaximin Novel Formulations in Patients with Sickle Cell Disease		Clinical Research Center (MASRI-CRC) 3-Zagazig University Hospital 4-Cairo University Hospital 5-Alexandria University Clinical Research Center			
64-	22/01/2023	AT/03A-017	Sponsor: Atea Pharmaceuti-cals CRO: Avicemer	A Phase 3 Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Efficacy and Safety of Bemnifosbuvir in High-Risk Outpatients with COVID-19	III	1- National Hepatology and Tropical Medicine Research Institute	Approved: 15/10/2023 Withdrawn 07/04/2024	COVID-19	(Pharmaceutical) Bemnifosbuvir/matching Placebo

Color Indicator	
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65-	13/02/2023	ENRICH-AF	Sponsor: Hamilton Health Science CRO: Clinmax	Edoxaban for Intracranial Haemorrhage Survivors with Atrial Fibrillation (ENRICH- AF)	IV	1-Ain Shams University Clinical Research Center (MASRI-CRC) 2-Zagazig University Hospital 3-Fayoum General Hospital 4-Tanta University Hospital 5-Mansoura University Hospital 6-Ain Shams Specialized Hospital 7-Alexandria University Clinical Research Center	Approved 10/05/2023 Recruitment Completion	Atrial Fibrillation in patients with previous Intracranial Haemorrhage	(Pharmaceutical) Edoxaban

Color Indicator	
Green	Biological
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Orange	Medical Device
Gray	Innovative
Red	Herbal

						8-Assuit University Hospital			
66-	13/02/2023	GBT440-038	Sponsor: GBT (Subsidiary of Pfizer)	An Open-Label Extension Study of Voxelotor Administered Orally to Paediatric Participants with Sickle Cell Disease Who Have Participated in Voxelotor Clinical Trials	III	1-Alexandria University Clinical Research Center 2- Zagazig University Hospital 3-Cairo University, Abu El Rich Hospital. 4- Ain Shams University, Faculty of Medicine CRC (MASRI).	Approved 30/03/2023 IMP Dosing Pause 02/05/2024 Early Terminated by the Sponsor 26/09/2024	Sickle Cell Disease	(Pharmaceutical) Voxelotor
67-	01/03/2023	GN41851 FENHANCE	Sponsor: Roche	A Phase III Multicentre, Randomized, Double-Blind, Double-Dummy,	III	1-Alexandria University-Clinical Research Center	Approved 26/4/2023 Withdrawn 11/01/2024	Relapsing multiple sclerosis	(Pharmaceutical) Fenebrutinib/ Teriflunomide/ matching placebo

Color Indicator	
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				Parallel-Group Study to Evaluate the Efficacy and Safety of Fenebrutinib Compared with Teriflunomide in Adult Patients with Relapsing Multiple Sclerosis. (FENHANCE)					
68-	06/03/2023	1305-0023 (FIBRONEER –ILD	Sponsor: Boehringer Ingelheim CRO: IQVIA	A Double Blind, Randomized, Placebo-Controlled Trial Evaluating the Efficacy and Safety of BI 1015550 Over At Least 52 Weeks in Patients with Progressive Fibrosing Interstitial Lung Diseases (PF-ILDs)	III	1-Ain Shams University Clinical Research Center (MASRI-CRC) 2- Alexandria University Clinical Research Center 3- Air Force Specialized Hospital	Approved 01/06/2023 Withdrawn 17/01/2024	Progressive Fibrosing Interstitial lung diseases (PF- ILDs)	(Pharmaceutical) BI 1015550 / matching placebo

Color Indicator	
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				((FIBRONEER – ILD)		4- Cairo University, Kasr Al Aini Hospital			
69-	06/03/2023	1305-0014 (FIBRONEER – IPF)	Sponsor: Boehringer Ingelheim CRO: IQVIA	A Double Blind, Randomized, Placebo-Controlled Trial Evaluating the Efficacy and Safety of BI 1015550 Over At Least 52 Weeks in Patients with Idiopathic Pulmonary Fibrosis (IPF) (FIBRONEER – IPF)	III	1- Ain Shams University Clinical Research Center (MASRI-CRC) 2- Alexandria University Clinical Research Center 3- Air Force Specialized Hospital 4- Cairo University, Kasr Al Ainy Hospital	Approved 01/06/2023 Withdrawn 08/01/2024	Idiopathic Pulmonary Fibrosis (IPF)	(Pharmaceutical) BI 1015550 / matching placebo
70-	16/03/2023	4202-HEM-201	Sponsor: Forma Therapeutics	A Phase 2 Open-Label Study to Evaluate Safety and Clinical Activity of FT-4202 in Patients	II	1- Cairo University, Abu El-Rich Children Hospital.	Approved 01/06/2023 Recruitment Completion	Thalassemia or Sickle Cell Disease	(Pharmaceutical) Etavopivat

Color Indicator	
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			CRO: MCT	with Thalassemia or Sickle Cell Disease		2-Cairo University, Kasr Al Eini Hospital.			
71-	15/05/2023	EFC16035 (PERSEUS)	Sponsor: Sanofi	A Phase 3, Randomized, Double-Blind, Efficacy and Safety Study Comparing SAR442168 to Placebo in Participants with Primary Progressive Multiple Sclerosis (PERSEUS)	III	1- Alexandria University Clinical Research Center	Approved 10/08/2023 Withdrawn 15/04/2024	Primary Progressive Multiple Sclerosis	(Pharmaceutical) Tolebrutinib/ Matching Placebo
72-	14/03/2024	WO43571 HereDERA	Sponsor: Roche	A Phase III, Randomized, Open-Label Study Evaluating the Efficacy and Safety of Giredestrant in Combination with	III	1- Sohag Oncology Center 2- Dar El Salam Cancer Hospital 3- National Cancer Institute	Approved 08/04/2024 Recruitment Completion	Previously Untreated Her2-Positive, Estrogen Receptor-Positive	Pharmaceutical Giredestrant

Color Indicator	
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				Phesgo Versus Phesgo After Induction Therapy with Phesgo+ Taxane in Patients with Previously Untreated Her2-Positive, Estrogen Receptor-Positive Locally-Advanced or Metastatic Breast Cancer (HereDERA)				Locally-Advanced or Metastatic Breast Cancer	
73-	22/04/2024	1517-CL-1003	Sponsor: Astellas Pharma Global Development CRO: MCT	A Phase 3, Open-label, Uncontrolled Study to Evaluate the Activity, Safety, Pharmacokinetics and Pharmacodynamics of Roxadustat for the Treatment of Anemia in	III	1- Cairo University Children's Hospital 2- Ain Shams University Hospital 3- Alexandria University Hospital	Approved 10/7/2024 Withdrawn 26/09/2024	Anemia in Pediatric Patients with Chronic Kidney Disease	Pharmaceutical Roxadustat

Color Indicator	
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				Pediatric Participants with chronic kidney disease 1517-CL-1003					
74-	05/06/2024	M23-698	Sponsor: Abbvie	A Phase 3, Randomized, Placebo-Controlled, Double-Blind Study to Evaluate Efficacy and Safety of Upadacitinib in Adult and Adolescent Subjects with Moderate to Severe Hidradenitis Suppurativa Who Have Failed Anti-TNF Therapy M23-698	III	1- Ain Shams University CRC (MASRI) 2- Air Force Specialized Hospital 3- Alexandria University CRC	Approved 07/08/2024 Withdrawn 09/04/2025	Moderate to Severe Hidradenitis Suppurativa	Pharmaceutical Upadacitinib
75-	16/12/2024	DAY101-002	Sponsor: Day One Biopharmaceuticals	A Phase 3, Randomized, International Multicenter Trial of DAY101	III	1- Children Cancer Hospital Egypt-57357	Approved 20/02/2025 Recruiting	Pediatric Low-Grade Glioma Harboring an Activating	Pharmaceutical Tovorafenib

Color Indicator	
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			CRO: MCT	Monotherapy Versus Standard of Care Chemotherapy in Patients with Pediatric Low-Grade Glioma Harboring an Activating RAF Alteration Requiring First-Line Systemic Therapy				RAF Alteration Requiring First-Line Systemic Therapy	
76-	29/01/2025	NN7535-7822 FLORAL	Sponsor: Forma Therapeutics CRO: IQVIA	An Open-Label, Multi-Centre, Rollover Study to Characterise Long-Term Safety and Efficacy of Etavopivat in Adults, Adolescents and Children Who Have Sickle Cell Disease or	III	1- Alexandria University Clinical Research Center (PI: Prof. Hoda Hassab) 2-Alexandria University Clinical Research Center (PI: Prof. Ashraf El Ghandour)	Approved 10/04/2025 Recruiting	Adults, Adolescents and Children Who Have Sickle Cell Disease or Thalassaemia	Pharmaceutical (Etavopivat)

Color Indicator	
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				Thalassaemia and Have Completed a Treatment Period in an Etavopivat Study (FLORAL)		3-Zagazig University Hospital 4-Kasr Al Aini Hospital, Cairo University 5-Ain Shams University Clinical Research Center (MASRI-CRC) 6- Abu El Rich Al Mounira Children Hospital, Cairo University			
77-	16/06/2025	ID-064A301 (OPUS-1)	Sponsor: Idorsia Pharmaceuticals CRO: RAY	A Phase 3, Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Study to Evaluate the Efficacy,	III	1- Air Force Specialized Hospital 2- Alexandria University Clinical Research Center 3- Ain Shams University	Approved 10/08/2025	Adult Subjects with Moderate to Severe Systemic Lupus Erythematosus	Pharmaceutical (Cenerimod)

Color Indicator	
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				Safety, and Tolerability of Cenerimod in Adult Subjects with Moderate-to-Severe Systemic Lupus Erythematosus (SLE) on Top of Background Therapy (OPUS-1)		Clinical Research Center (MASRI-CRC)			
78-	20/07/2025	MK4482-0023	Sponsor: MSD	A Phase 3, Randomized, Placebo-Controlled, Double-Blind Clinical Study to Evaluate the Efficacy and Safety of Molnupiravir (MK-4482) in Non-Hospitalized Adults With COVID-19 at High Risk for	III	1- Air Force Specialized Hospital	Submission Cancellation by EDA 04/11/2025	Non-Hospitalized Adults With COVID-19 at High Risk for Disease Progression	Pharmaceutical (Molnupiravir)

Color Indicator	
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				Disease Progression					
79-	31/07/2025	1397-0014	Sponsor: Boehringer Ingelheim International GmbH CRO: IQVIA	A Phase III, Randomized, Double-Blind, Placebo-Controlled Study to Assess the Efficacy, Safety, and Tolerability of BI 1291583 2.5 mg Administered Once Daily for Up to 76 Weeks in Patients With Bronchiectasis (The AIRTIVITY® Study)	III	**1-** Air Force Specialized Hospital **2-** Alexandria CRC, Faculty of Medicine Alexandria **3-** Kasr Al Aini Hospital, Cairo University, 4- MASRI CRC, Faculty of Medicine, Ain Shams University,	Approved 08/10/2025	Bronchiectasis	Pharmaceutical (BI 1291583 2.5 mg)
80-	25/08/2025	D6972C00002	Sponsor: Astra Zeneca	A Phase III, Randomised, Double-blind, Placebo-controlled, Event-driven Study	III	1- Alexandria University Clinical Research Center,	Approved 11/11/2025	Chronic Kidney Disease and High	Pharmaceutical (Baxdrostat & Dapagliflozin)

Color Indicator	
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			CRO: IQVIA	to Assess the Efficacy, Safety and Tolerability of Baxdrostat in Combination with Dapagliflozin Compared with Dapagliflozin Alone on Renal Outcomes and Cardiovascular Mortality in Participants with Chronic Kidney Disease and High Blood Pressure (Bax-Duo PACIFIC)		2- Ain Shams University, MASRI Clinical Research Center 3- National Institute of Urology and Nephrology, 4- Mansoura University, Urology & Nephrology Center, 5- Mansoura University, Specialized Medical Hospital,		Blood Pressure	
81-	08/09/2025	1378-0041	Sponsor: Boehringer Ingelheim	A Phase III Double-Blind, Randomised, Parallel-Group Superiority	III	1- Alexandria University Hospital,	Withdrawn 10/12/2025	Type 2 Diabetes, Hypertension and Established	Pharmaceutical (Vicadrostat (BI 690517) and Empagliflozin)

Color Indicator	
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			International GmbH CRO: IQVIA	Trial to Evaluate Efficacy and Safety of the Combined Use of Oral Vicadrostat (BI 690517) and Empagliflozin Compared With Placebo and Empagliflozin in Participants with Type 2 Diabetes, Hypertension and Established Cardiovascular Disease		2- Ain Shams University Hospital, 3- Cairo University Hospital,		Cardiovascular Disease	
82-	18/09/2025	AT-01B-008	Sponsor: Atea Pharmaceuticals CRO: AVICEMER	An Evaluation of Bemnifosbuvir-Ruzasvir (BEM/RZR) Versus Sofosbuvir-Velpatasvir (SOF/VEL) for the Treatment of	III	1.National Hepatology and Tropical Medicine Institute, Clinical Research Center	Approved 15/12/2025	Chronic Hepatitis C Virus (HCV)	Pharmaceutical (Bemnifosbuvir-Ruzasvir)

Color Indicator	
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				Chronic Hepatitis C Virus (HCV) Infection in a Phase 3 Randomized, Controlled, Open-label Study		2.Air Force Specialized Hospital Research Center			
83-	18/09/2025	GLOBOTRK	Sponsor: St. Jude Children's Research Hospital CRO: MCT	Phase 2 Study of Entrectinib as a Single Agent in Upfront Therapy for Children <3 Years of Age with NTRK1/2/3 or ROS-1-Fused CNS Tumors (GLOBOTRK)	II	Children Cancer Hospital Egypt 57357	Approved 18/12/2025	NTRK1/2/3 or ROS-1-Fused CNS Tumors	Pharmaceutical (Entrectinib)
84-	24/07/2022	MD-004	Sponsor: Ezz Medical Industries CRO: Dataclin	Open labelled non randomized self-controlled study to evaluate the safety and performance of Ezvent in hospitalized	III (pivotal)	1-Kasr Al-Aini university Hospital	Approved 28/08/2022 Suspended 01/01/2024 Resumed	Hospitalized mechanically ventilated patients	Medical device (Ezvent)

Color Indicator	
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Red	Herbal

				mechanically ventilated patients			13/01/2024 Completed 03/06/2025		
85-	15/05/2022	COAV101 B12301	Sponsor: Novartis CRO: MCT	A randomized sham controlled double –blind study to evaluate the efficacy and safety of intrathecal (IT) QAV101 in patients with later onset type 2 spinal muscular atrophy (SMA) who are ≥2 to <18 years of age, treatment naïve sitting and never ambulatory	III	1-Department of Neurology, Ain Shams University Specialized Hospital.	Approved 02/08/2022 Early terminated (by sponsor) 18/12/2023	type 2 spinal muscular atrophy (SMA)	Innovative QAV101 (Zolgensma) (Onasemnogene abeparvovec)
86-	06/06/2023	Urso-003	Sponsor: Minapharm	Multi-Center, randomized, control, phase IV trial to compare the efficacy & safety	IV	1-Clinical Research Center, Air force	Approved 18/09/2023 Suspended	Compensated Chronic Liver Disease Patients	Innovative Ursoplus® capsules/

Color Indicator	
Green	Biological
Blue	Pharmaceutical
Orange	Medical Device
Gray	Innovative
Red	Herbal

			CRO: Dataclin	of Ursoplus® capsules (UDCA 250mg & Silymarin 140mg) versus UDCA alone versus Placebo among Compensated Chronic Liver Disease Patients		specialized Hospital 2-National Hepatology and Tropical Research Institute (NHTMRI)	(Recruitment suspension) 26/11/2024		Ursofalk® capsules
87-	06/06/2023	Cipro-001	Sponsor: Minapharm, CRO: Nagy Research	Single center, Open Label, controlled Study to assess the safety & efficacy of Oral Ciprodiazole ® Tablets (Ciprofolxacin/ Metronidazole) versus currently used Ciprofloxacin Tablets & Metronidazole tablets in pelvi-abdominal	IV	1- General Syrgery department, Menoufia University Hospital.	Suspended 12/09/2023	Pelvi-abdominal infections and following IV antibiotics in post-operative period, for pelvi-abdominal surgeries or acute conditions	Innovative Ciprodiazole ® Tablets (Ciprofolxacin/ Metronidazole)

Color Indicator	
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Orange	Medical Device
Gray	Innovative
Red	Herbal

				infections and following IV antibiotics in post-operative period, for pelvi-abdominal surgeries or acute conditions					
88-	15/05/2023	Sub-Thromb-001	Sponsor: Minapharm	A Prospective, Single- Center, Phase IV Interventional, Single Arm Trial for the Evaluation of subcutaneous recombinant Hirudin 15 mg (RB variant) in prophylaxis of Deep Vein Thrombosis (DVT) post major orthopedic operations	IV	1- Department of Orthopedics and Trauma Surgery, El-Hadra University Hospital	Withdrawn 28/08/2023	prophylaxis of Deep Vein Thrombosis (DVT) post major orthopedic operations	Innovative Thrombex (recombinant Hirudin)

Color Indicator	
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Orange	Medical Device
Gray	Innovative
Red	Herbal

89-	24/10/2023	GRC/NE-CV/EG/39 /IV	Sponsor: Nerhadou Internatio-nal CRO: Genuine research center	A prospective, Multicentre, Open-label, Single-arm Interventional Study of Bisoprolol (Nerkardou) (Between Low Dose and High Dose) 5 and 10 mg ODF Treatment in Egyptian Patients with Essential Hypertension	IV	1- Department of General Internal Medicine , Beni-Suef University Hospital 2- Department of Cardiology and vascular medicine, Fayoum University Hospital	Approved 10-3-2024 Terminated by sponsor 13/05/2025	Essential Hypertension	Innovative Nerkardou (Bisoprolol) Oral dispersible film

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Gray	Innovative
Red	Herbal