

## Non Government Porject

| Sr.<br>No | Name of the Project, Clinical Trial,<br>Endowment, Chairs  | Name of the Principal<br>Investigator/ Co<br>Investigator | Name of the funding<br>agency            | Type<br>(Govermentt/<br>Non-Goverment | Department of<br>Pricipal<br>Investigator/<br>Co- Inveatigator | Year of<br>Award | Funds<br>provided<br>(INR in<br>Lakhs) | Duration of<br>Project |  |
|-----------|--|---|--|---------------------------------------|--|------------------|--|------------------------|--|
|           |  |   | 2014-15                                  |                                       |  |                  |  |                        |  |
| 1         | Validation of ingenious device designed by<br>ACTOFIT for kinematic analysis of joint motion   | Dr. Rajani Mullerpatan                                    | ACTOFIT                                  | Non - Goverment                       | Physiotherapy Navi<br>Mumbai                                   | 2014             | 0.3                                    | 1 yrs                  |  |
| 2         | Establishment of MGM Centre of Human<br>Movement Science at MGM School of<br>Physiotherapy   | Dr. Rajani Mullerpatan                                    | International Society of<br>Biomechanics | Non - Goverment                       | Physiotherapy Navi<br>Mumbai                                   | 2014             | 74.6683                                | 3 yrs                  |  |
|           | 2015-16  |   |  |                                       |  |                  |  |                        |  |
| 1         | Non Invasive TB Triage & Patient Mapping<br>platform using breath via Low cost titanium<br>dioxide Nanotube sensor   | Dr. Samir Pachpute (PI)                                   | IKP Knowledge Park                       | Non - Goverment                       | Microbiology Navi<br>Mumbai                                    | 2015             | 49.9                                   | 1 yrs                  |  |
| 2         | The effects of labour and birth positioning on<br>pelvic dimensions: gaining further insight to<br>improve the birth experience  | Dr. Rajani Mullerpatan                                    | Shastri foundation                       | Non - Goverment                       | Physiotherapy Navi<br>Mumbai                                   | 2015             | 1.12                                   | 1 yrs                  |  |
| 3         | Clinical Evaluation of Breath-Based point of<br>care test for diagnosis of Pulmonary<br>Tublerculosis  | Dr. Pradeep Potdar  | University of UTAH,<br>USA & Nanosynth   | Non - Goverment                       | Respiratory<br>Medicine  | 2016             | 66                                     | 1 yrs                  |  |
| 4         | A Randomized, Double-blind, Parallel Group<br>Study to Evaluate the Effect and Safety of Test<br>Biscuits as Compared to Placebo Biscuits in<br>Regulating Blood Glucose Level in Subjects<br>With Type 2 Diabetes Mellitus. | Dr. Deepak Bhosle   | Karmik life sciences                     | Non - Goverment                       | Pharmacology<br>Aurangabad                                     | 2016             | 3.27866                                | Ongoing                |  |

| 5 | A Randomized, Double-Blind, Placebo-<br>Controlled, Three-Arm, Parallel Group, Multi-<br>Centric, Clinical Study To Evaluate The<br>Therapeutic Bio-Equivalence Of Two<br>Tacrolimus 0.1% Topical Ointment<br>Formulations In Adult Patients With Moderate<br>To Severe Atopic Dermatitis.  | Dr. Ashish Deshmukh       | Lambda Therapeutic<br>Research Limited                 | Non - Goverment | TB & Chest<br>Aurangabad   | 2016 | 0.9576  | Ongoing |
|---|---|---------------------------|--|-----------------|----------------------------|------|---------|---------|
| 6 | Arandomized, multi center, open label, two-<br>treatment, two-period, two-sequence, multiple<br>dose,crossover, steady state bioequivalence<br>study of Everolimus tablets, 10 mg of Biocon<br>Limited, India vs. Afinitor@ (Everolimus)<br>tablets, 10 mg of Novartis Phannaceuticals<br>Corporation, USA in<br>advanced renal cell carcinoma (RCC) patients | Dr. Chandrashekhar Tamane | Veeda Clinical Research<br>Pvt. Ltd,                   | Non - Goverment | Oncology<br>Aurangabad     | 2016 | 1.12928 | Ongoing |
|   |   |                           | 2016-17  |                 |                            |      |         |         |
|   | A Randomized, Open Label, Two Period, Single<br>Dose, Crossover, Bioavailability Study Of<br>Paclitaxel Injection Concentrate For Nano-<br>Dispersion (PICN) And Abraxane® In Subjects<br>With Locally Recurrent Or Metastatic Breast<br>Cancer.  | Dr. Chandrashekhar Tamane | Sun Pharma Advanced<br>Research Company Ltd<br>(SPARC) | Non - Goverment | Oncology<br>Aurangabad     | 2016 | 1.12928 | Ongoing |
| 2 | Real world, Non-interventional, Observational<br>Study of Hydroxyzine hydrochloride in chronic<br>pruritus  | Dr. H R Jerajani          | Clinresearch Healthcare<br>solution pvt ltd            | Non - Goverment | Dermatology Navi<br>Mumbai | 2016 | 2.1     | Ongoing |
|   | A Phase IV, Open-Label, Multi-center Study to<br>Evaluate the Safety of Apixaban in Indian<br>Subjects Undergoing Elective Total Knee<br>Replacement or Total Hip Replacement Surgery.<br>Protocol Number : CV185-158   | Dr. Girish Gadekar        | PPD Pharmaceutical<br>Development India Pvt<br>Ltd     | Non - Goverment | Orthopaedics<br>Aurangabad | 2016 | 19.2    | Ongoing |
| 4 | Real world, non-interventional, observational<br>study of Venusia Max Cream as Moisturizer in<br>Psoriasis  | Dr. H R Jerajani          | Bio Quest pvt ltd                                      | Non - Goverment | Dermatology Navi<br>Mumbai | 2017 | 2.1     | Ongoing |
| 5 | To evaluate efficacy and safety of Bacillus<br>clausii (2 billion spores/5 ml) suspension as an<br>add on therapy to standard of care in acute viral<br>diarrhoea in children   | Dr. Nimain Mohanty        | Wockhardt Ltd  | Non - Goverment | Paediatrics Navi<br>Mumbai | 2017 | 11.49   | Ongoing |

| 6  | A Multicenter, Randomized, Double-Blind,<br>Vehicle-Controlled Phase II Study to Evaluate<br>the Efficacy, Tolerability, and Safety of Topical<br>Povidone-Iodine (PVP-I, 2% [W/W]) in<br>Pediatric Subjects for the Treatment of<br>MolluscumContagiosum.   | Dr. Ashish Deshmukh   | JSS Clinical Research<br>Limited | Non - Goverment | TB & Chest<br>Aurangabad | 2017 | 3.495  | Ongoing |
|----|--|-----------------------|----------------------------------|-----------------|--------------------------|------|--------|---------|
| 7  | A Randomized, Double-Blind, Placebo-<br>Controlled, Phase 2 Study to Assess the<br>Efficacy, Pharmacokinetics,<br>Pharmacodynamics and Safety of LNP1892<br>(Monotherapy) in Chronic Kidney Disease<br>(CKD) Patients with Secondary<br>Hyperparathyroidism (SHPT), On Dialysis and<br>Not on Dialysis. Protocol Number:<br>LRP/LNP1892/2016/007; Version No.: 1.2,<br>Dated 15 Dec 2016 | Dr. Sudhir Kulkarni   | LUPIN                            | Non - Goverment | Nephrology<br>Aurangabad | 2017 | 1      | Ongoing |
| 8  | A rarulomized, open label, patallel-group, actjle-<br>comparator controlled, muhi-center study to<br>eraludte tfu efrcacy Ind safety ofLlipristal<br>acetate (5 ng tablets), as compared with<br>Leuprolide acetate (3.75 mg intramusculal<br>injection) for 12 weeks, in the preoryratiye<br>h,eatment of moderate to severe swptomatic<br>uterine<br>fibroids                          | Dr. Lakshmi Rachkonda | Cliantha Research<br>Limited     | Non - Goverment | OBGY Aurangabad          | 2017 | 0.7902 | Ongoing |
| 9  | Title An open label, Two Arms, Comparaiive,<br>Phase'IV Clinical Study evaluatLng safely and<br>efficacy of Oratil LZ (combination of<br>Cefuroxime 250m9 + Linezo id 600m9) versus L<br>nezolid 600m9 in patiefts with Diabetic Foot<br>Infectons.  | Dr. Anuradha Patil    | Macleods<br>Pharmaceutical Ltd   | Non - Goverment | Pathology                | 2017 | 1.4375 | Ongoing |
| 10 | An Open Label Prospective, comparative,<br>randomized, clinical study evaluating efficacy<br>and safety of treatment A (Tamsulocin 0.4 mg<br>Modified release capsule) versus treatment B<br>(FDC Deflazacort 30 mg plus Tamsulocin 0.4<br>mg Tablet) in the patients with ureteral stone.   | Dr. Prashant Darakh   | Macleods<br>Pharmaceutical Ltd   | Non - Goverment | Urology<br>Aurangabad    | 2017 | 2.4    | Ongoing |

| 11 | A Two arm, Comparative, Parallel, Randomized,<br>double-blind, double-dummy Clinical study to<br>evaluate and compare the efficacy and safety of<br>Cefrine (combination of cefdinir 300 mg +<br>lactobacillus 60 million cells) Capsule versus<br>Cefuroxime 250 mg Tablet in the treatment of<br>patients with Upper Respiratory Tract<br>Infections.    | Dr. Rajendra Bohra  | Macleods<br>Pharmaceutical Ltd  | Non - Goverment | ENT Aurangabad             | 2017 | 2.4     | Ongoing |
|----|--|---------------------|---------------------------------|-----------------|----------------------------|------|---------|---------|
| 12 | A ProsPective, I\rulticentric, Phase IV Clinical<br>Study evaluating safety and efficacy of<br>Leuprorelin 3.5mg InjectIon plus Enzomac<br>Tablet (Trypsin 96 mg + Bromelain 180 mg +<br>Rutoside Trihydrate 200 mg) versus Leuproretin<br>3 smg Injection in the treatment of patients<br>diagnosiswth Endometrio6is<br>Protocol No.: CT-205-LEEN-2016    | Dr. Swati Shiradkar | Macleods<br>Pharmaceutical Ltd  | Non - Goverment | Community<br>Medicine      | 2017 | 1.725   | Ongoing |
| 13 | An open label, Two arm, Comparative,<br>Randomized Phase IV Clinical Study evaluating<br>efficacy and safety of Alrista Forte (Epalrestat<br>150 Mg + Methylcobalmin 1500 Mcg +<br>Pregabalin 150 Mg) Tablet versus<br>Pregabalin 150 Mg Capsule in patients with<br>Diabetic Neuropathy   | Dr. Deepak Bhosle   | Macleods<br>Pharmaceutical Ltd  | Non - Goverment | Pharmacology<br>Aurangabad | 2017 | 2.4     | Ongoing |
|    |  |                     | 2017-18                         |                 |                            | •    |         |         |
| 1  | 26 Week, Multicenter, randomized, placcbo<br>controlled, double blind, parallel group, phase 3<br>trial with a 26 week safety extension period<br>evaluting the safety and efficacy of<br>Dapagliflozin 5 and 10 mg and saxagliptin 2.5<br>and 5 mg in pediatric patients with type 2<br>diabetes mellitus who are between 10 and below<br>18 years of age | Dr. Deepak Bhosale  | Astrazeneca Pharma<br>India Ltd | Non - Goverment | Pharmacology<br>Aurangabad | 2017 | 3.77323 | Ongoing |

| 2 | A 24'week, randomised, double'blind, double-<br>dummv parallel group, multi cenire,<br>active'controlled study to evaluate cfficacy and<br>safety ofremogliflozin etabonate in subjects with<br>type'2 diabetes mellitus.  | Dr. Deepak Bhosale       | Glenmark<br>Pharmaceutical Ltd                           | Non - Goverment | Pharmacology<br>Aurangabad    | 2017 | 25.65    | Ongoing |
|---|--|--------------------------|--|-----------------|-------------------------------|------|----------|---------|
| 3 | A Study to Evaluate the Effect of Dapagliflozin<br>on Incidence of Worsening Hearl Failure or<br>Cardiovascular Death in in Patients lvith phronic<br>Heart Failure with Reduced Ejection F-raction.   | Dr. Prashant Udgire      | Astrazeneka Pharma<br>India Ltd                          | Non - Goverment | Plastic Surgery<br>Aurangabad | 2017 | 63.3     | Ongoing |
| 4 | Inj- TR2SERVE :A comprehensive program to<br>train, triage, and improve services for<br>unintended childhood injuries in India   | Dr. Maninder singh Setia | Grand Challenges<br>Canada                               | Non - Goverment | OBGY Navi<br>Mumbai           | 2018 | 0.98545  | Ongoing |
| 5 | Phase 3 Multicenter Double blind study to<br>evaluate the long term safety and efficacy of<br>barciunib in adult patients with atopic dermatitis   | Dr. H R Jerajani         | Elli Lilly Company<br>(India) Pvt Ltd Gurgaon<br>Haryana | Non - Goverment | Dermatology Navi<br>Mumbai    | 2018 | 24.25868 | Ongoing |
| 6 | A Multicenter Randomized double blind placebo<br>controlled study to evaluate the efficacy and<br>safety of baricitinib in adult patients with<br>moderate to severe atopic dermatitis   | Dr H R Jerajani          | Elli Lilly Company<br>(India) Pvt Ltd Gurgaon<br>Haryana | Non - Goverment | Dermatology Navi<br>Mumbai    | 2018 | 15.7825  | Ongoing |
|   |  | L                        | 2018 - 19  | L               |                               |      |          |         |
| 1 | Biomechanical analysis and energy expenditure<br>of traditional, chair and wall Suryanamaskar  | Dr. Rajani Mullerpatan   | Sancheti College of<br>Physiotherapy                     | Non - Goverment | Physiotherapy Navi<br>Mumbai  | 2018 | 0.172    | 1 yrs   |
| 2 | A Randomized, Double-blind, Placebo-<br>controlled, Parallel-group, Multicenter Study to<br>Demonstrate the Effects of Sotagliflozin on<br>Cardiovascular and Renal Events in Patients<br>with Type 2 Diabetes, Cardiovascular Risk<br>Factors and Moderately Impaired Renal<br>Function | Dr. Prashant Udgire      | Sanofi- Synthelabo Pvt<br>Ltd                            | Non - Goverment | Cardiology<br>Aurangabad      | 2018 | 112.925  | Ongoing |

| 3 | A Randomized, Double-Blind, Placebo-<br>Controlled, three-arm, Parallel Design, Multiple<br>site, Study to Evaluate the Therapeutic<br>equivalence and Safety of Tacrolimus<br>Ointment,0.1% (Encube Ethicals Private<br>Limited) with Protopic® - (Tacrolimus<br>Ointment 0.1% (Astellas Pharma US, Inc) in the<br>treatment of Moderate to Severe Atopic<br>Dermatitis. | Dr. Ashish Deshmukh               | Accutest  | Non - Goverment | Skin and VD<br>Aurangabad                 | 2018 | 10.97                          | Ongoing  |
|---|---|-----------------------------------|---|-----------------|---|------|--------------------------------|----------|
| 4 | A Multicentric, Open lable, Randomized,<br>Comparative, clinical study evaluating safety<br>and Efficacy of Fixed Dose combination of<br>Trypsin 48 mg + Bromelain 90 mg + Rutoside<br>Trihydrate 100 mg enteric coated tablet versus<br>Serratiopeptide 10 mg enteric coated tablet in<br>patient for heling potential in surgical wound<br>after minor surgery.         | Dr. Mahendra Surywanshi           | Macleods<br>Pharmaceutical Ltd  | Non - Goverment | Surgery<br>Aurangabad                     | 2018 | 2.88                           | Ongoing  |
| 5 | A Phase III Randomized, Double Blind, Parallel<br>Group, Placebo Controlled, Multi-Centre,<br>Multinational Study to Evaluate Efficacy and<br>Safety of TRC150094 as an Add on to Standard<br>of Care in Improving Cardiovascular Risk in<br>Subjects with Diabetes, Dyslipidemia and<br>Hypertension.  | Dr. Deepak Bhosle                 | IQVIA HDS (India) Priva   | Non - Goverment | Pharmacology<br>Aurangabad                | 2019 | 20.3976                        | Ongoing  |
| 6 | Safety and Efficacy of Lipiodol® Ultra<br>Fluid in Association with Surgical Glues during<br>Vascular Embolization,<br>a phase IV study.  | Dr.Shivaji Pole                   | SIRO Clinpharm<br>Pvt. Ltd.   | Non - Goverment | Interventional<br>Radiology<br>Aurangabad | 2019 | 10.7085                        | Ongoing  |
|   |   |                                   | 2019 -2020  |                 |   |      |                                |          |
| 1 | Rotasili ® vaccine intussusceptions surveillance<br>in Kerala, Karnataka, Maharashtra and<br>Gurjarat, India  | Dr. Mohd Haseeb                   | Bill and Melinda Gates<br>Foundation                                      | Non- Goverment  | Medicine                                  | 2020 | 383400                         | 2 year   |
| 2 | Major neurological complications following<br>central neural blockade – A pilot study in<br>Aurangabad city   | Dr. Sadhana Kulkarni              | MGM Institute of<br>Health Sciences, Navi<br>Mumbai (faculty project<br>) | Non- Goverment  | Emergency<br>Medicine                     | 2019 | 25,000                         | 1 Year   |
| 3 | Safety and Efficacy of Lipiodol Ultra fluid in<br>association with Surgical Glues during Vascular<br>Embolization. (IV) Protocol No: LUF-44-001   | Dr Shivaji Pole,<br>Dr Ejaj Patel | GUERBET   | Non- Goverment  | Radiology                                 | 2019 | Rs.1,07,085/-<br>(Per Patient) | 180 days |

| 4 | A Multicentric, Randomized, Open label,<br>Clinical trial To Evaluate Efficacy and Safety of<br>Tapentadol Nasal Spray in comparison to<br>Tramadol Immediate release Capsule and<br>Intravenous injection in patients with Post<br>operative moderate to severe pain.<br>PNo: CT/TAPE/PAIN/17/03  | Dr Tejendersingh Chhabda,<br>Dr Amey Chakkarwar | SIRO CRO                         | Non- Goverment | Surgery         | 2019 | Rs. 27,600/-<br>(Per Patient) | 180 days |
|---|--|---|----------------------------------|----------------|-----------------|------|-------------------------------|----------|
| 5 | Multi centre cross sectional epidemiological<br>study to characterize the prelevance and<br>distribution of lipoprotein (a) levels among<br>patient with established cardiovascular<br>diseaseP No : CTQJ23A12001  | Dr Prashant Udgire                              | Novartis                         | Non- Goverment | Cardiology      | 2019 | Rs 8,400/-<br>(Per Patient)   | 180 days |
| 6 | A Randomized,Open label,<br>Prospective,Comparative Multicentre, Parallel<br>group Active controlled Phase III Study<br>evaluate safety and efficacy of Ropivacaine<br>readyfusor 2mg/ml versus Ropivacaine ballon<br>pump infusor 2mg/ml as continuous surgical<br>site infusion for treatment of post surgical Pain<br>in lower abdominal Laprotomy.<br>P. No:CP/04/18 | Dr Vasanti Kelkar                               | JSS                              | Non- Goverment | Anaesthesiology | 2019 | Rs 8,400/-<br>(Per Patient)   | 180 days |
| 7 | A Prospective, Single-center, Open-label,<br>Randomized, Parallel group, Active-controlled<br>Clinical Study to Evaluate the Efficacy and<br>Safety of CefpodoximeProxetil 200 mg +<br>Clavulanic acid 125 mg versus Amoxicillin 500<br>mg + Clavulanic acid 125 mg in the Treatment<br>of Patients with Upper Respiratory Tract<br>Infections.                          | Dr Rajendra Bohra                               | Macleods<br>Pharmaceuticals Ltd. | Non- Goverment | ENT             | 2019 | Rs 6,000/-<br>per patient     | 180 days |

| 8  | A Prospective, ,Double Blind, Randomized<br>Parallelgroup,vehicle controlled,Multicentre<br>Clinical Study to evaluate Safety and<br>Bioequivalence of Mupirocin Cream USP 2%<br>(Supplied by Glasshouse Pharmaceuticals<br>Limited Canada) using Mupirocin cream 2%<br>(Manufactured by :Glenmark Pharmaceuticals<br>Inc:USA) as a reference product,in subjects with<br>secondarily Infected Traumatic Skin Lesions<br>P.No : GH-SL-IN-01 | Dr.Ashish Deshmukh   | JSS                                     | Non- Goverment | Dermatology  | 2019 | Rs 15,000 per<br>patient   | 180 days |
|----|---|--|---|----------------|--|------|----------------------------|----------|
| 9  | A randomized, double-blind, placebo-<br>controlled, study evaluating the efficacy and<br>safety of otilimab IV in patients with severe<br>pulmonary COVID-19 related disease  | Dr Umar Quadri   | GSK/PPD                                 | Non- Goverment | Emergency<br>Medicine  | 2020 | Rs 7,27,291/-              | 1 Year   |
| 10 | A Phase II, controlled clinical study designed to<br>evaluate the effect of ArtemiC in patients<br>diagnosed with COVID-19.   | Dr. Umar Quadri  | Biosphere Clinical<br>Research Pvt. Ltd | Non- Goverment | Emergency<br>Medicine  | 2020 | Rs 18,000/-                | 1 Year   |
| 11 | A prospective, multi-centre, open label, phase<br>IV study to evaluate safety and efficacy profile<br>of Infimab <sup>™</sup> in patients with moderate to<br>severe plaque psoriasis   | Dr. Ashish Deshmukh  | Reliance Life Sciences                  | Non- Goverment | Dermatology &<br>Pharmacology  | 2020 | Rs 36,344/-                | 1 Year   |
| 12 | An open label, randomized, active-controlled,<br>multi-centric phase-ii/iii study in indian<br>toddlers and infants to assess the<br>immunogenicity and safety of siiplhexasiiltm<br>(dtwp-hepb-ipv-hib) vaccine in comparison<br>with siiplpentavac (dtwp-hepb-hib) + poliovac<br>(ipv) vaccines, administered as separate<br>injections.  | Dr. Deepak Tayade  | Serum Institute of India                | Non- Goverment | Community<br>Medicine  | 2020 | Rs 26,300/-                | 1 Year   |
| 13 | A randomized double blind placebo controlled<br>study to evaluate the effcacy and safety of<br>cordycepes capsules (Food supplment) as an<br>additional on therapy in patient with mild to  | Dr. Sagar Sinha<br>Dr. Siddharth Dubhasi,<br>(AIIMS, Nagpur)<br>Dr. Jayashree Ghanekar,<br>Dr. Sameer Kadam,<br>Dr. Parineeta Samant | M/S Ambosia Food<br>Farm Co. Uttrakhand | Non- Goverment | Emergecny<br>Medicine , Surgery,<br>Biochemistry<br>and AIIMS,<br>Nagpur | 2020 | 14,72,000                  | 1 Year   |
| 14 | Survey to collect voice samples of patients and<br>age matched healthy individuals P No:<br>VoQuest01   | Dr. Deepak Bhosle  | Quest                                   | Non- Goverment | Pharmacology   | 2020 | Rs.100 /-<br>(Per Patient) | 180 days |

| 15 | A Multicenter, Phase III, Double-Blind,<br>Randomized, Placebocontrolled Study To<br>Evaluate The Efficacy Of Recombinant Bcg<br>Vpm1002 In Reducing Infection Incidence And<br>Disease Severity Of Sars-Cov-2/Covid-19<br>Among High-Risk Subjects P No: SII-<br>Rbcg/COVID-19/IN-01   | Dr. Deepak Tayade        | Serum Institute of India<br>Pvt. Ltd | Non- Goverment | Community<br>Medicine | 2020 | Rs 18,200/-<br>(Per Patient) | 180 days |
|----|---|--------------------------|--------------------------------------|----------------|-----------------------|------|------------------------------|----------|
|    | A Phase III,Multicentre, Randomized, Double<br>Blind, Parallel group,Comparative Clinical<br>Study to evaluate Efficacy and Safety of<br>Ropivacaine Hydrochloride 0.75% (7.5mg/ml)<br>in Dextrose 8% (80 mg/ml) injection compared<br>to Bupivacaine Hydrochloride 0.5% (5mg/ml)<br>in Dextrose 8% (80mg/ml) injection for subjects<br>undergoing lower limb orthopedic surgeries<br>under spinal anaesthesia P No: BCR/RPD/01 | Dr.Vasanti Kelkar        | Neon                                 | Non- Goverment | Anaesthesiology       | 2020 | RS 10,000/-<br>(Per Patient) | 180 days |
| 17 | A Comparative,Randomized,Two arm, Double<br>blind,Parallel Group,Multicentre, Phase III<br>Clinical study to Evaluate the<br>Efficacy,Safety,and Tolerability of Netarsudil<br>Opthalmic Solution 0.02% vs Timolol maleate<br>Eye drops 0.5% in treatment of elevated<br>Intraocular Pressure(IOP) in subjects with Open<br>Angle Glaucoma or Ocular Hypertension P No:<br>APL/CT/18/03   | Dr. Jyotika Mishrikotkar | Mediclin Clinical<br>Research        | Non- Goverment | Ophthalmology         | 2020 | Rs 18,500 (Per<br>Patient)   | 180 days |
| 18 | A prospective, Multicentre, Randomized,<br>Double blind ,Phase3 study to evaluate the<br>safety and efficacy of a gel formulation of<br>Esmolol hydrochloride (Galnobax) in treating<br>Diabetic Foot Ulcer<br>P No: NG-A16   | Deepak Bhosale           | JSS                                  | Non- Goverment | Pharmacology          | 2020 | Rs 80,000 (Per<br>Patient)   | 180 days |
| 19 | A Randomized Open-Label Study To Evaluate<br>The Efficacy And Safety Of Favipiravir And<br>Umifenovir As Compared To Favipiravir Alone<br>In Moderate Hospitalized Adult Indian Covid-<br>19 Patients P No: GPL/CT/2020/004/III   | Dr. Anand Nikalje        | Glenmark                             | Non- Goverment | Pharmacology          | 2020 | Rs 95,000/-<br>(Per Patient) | 180 days |
| 20 | A Phase II, controlled clinical study designed to<br>evaluate the effect of ArtemiC in patients<br>diagnosed with COVID-19.   | Dr. Umar Quadri          | MGC Pharma                           | Non- Goverment | Emergency<br>Medicine | 2020 | Rs 18,000/-<br>per patient   | 180 days |

| 21 | A Randomized, Open Label, Single Centre,<br>Observational, Prospective, Clinical Study to<br>Evaluate the Efficacy and Safety of GanjhuVir<br>syrup and tablet when administered along with<br>Standard of Care (SOC) and compared against<br>SOC in Covid-19 positive patients.  | Dr.Umar Quadri       | Radhika Ayurveda<br>Research and<br>Development   | Non- Goverment | Emergency<br>Medicine | 2020 | Rs 6,666/-<br>per patient    | 180 days |
|----|---|----------------------|---|----------------|-----------------------|------|------------------------------|----------|
| 22 | A Randomized, Open Label, Parallel Efficacy,<br>Active Control, Multi- Centre Exploratory Drug<br>Trial to Evaluate Efficacy and Safety of an<br>Ayurvedic Formulation- II (SanshamaniVati<br>Plus) as Adjunct Treatment to Standard of Care<br>for the management of Mild to Moderate<br>COVID-19 Patients.                              | Dr. Syed Umar Qudari | AYUSH CSIR  | Non- Goverment | Emergency<br>Medicine | 2020 | Rs 15,000/-<br>per patient   | 180 days |
| 23 | An open label, randomized, active-controlled,<br>multi-centric phase-ii/iii study in indian<br>toddlers and infants to assess the<br>immunogenicity and safety of siiplhexasiiltm<br>(dtwp-hepb-ipv-hib) vaccine in comparison<br>with siiplpentavac (dtwp-hepb-hib) + poliovac<br>(ipv) vaccines, administered as separate<br>injections | Dr. Deepak Tayade    | Serum Institute of India                          | Non- Goverment | Community<br>Medicine | 2020 | Rs 26,300/-<br>per patient   | 180 days |
| 24 | A randomized, double-blind, placebo-<br>controlled, study evaluating the efficacy and<br>safety of otilimab IV in patients with severe<br>pulmonary COVID-19 related disease  | Dr. Umar Quadri      | GSK   | Non- Goverment | Emergency<br>Medicine | 2020 | Rs 7,27,291/-<br>per patient | 180 days |
|    |   |                      | 2020- 2021  |                |                       |      |                              |          |
| 1  | Randomized, double-blind, placebo-controlled,<br>parallel-group, multi-centre phaseII/III<br>adaptive clinical trial to assess the safety and<br>immunogenicity, of Gam-COVID-Vac<br>cornbinedvector viiccine for SARS-CoV-2<br>infection in Indian Healthy Subjects.   | Dr. Anand Nikalje    | JSS Medical Research<br>India Pvt Ltd.,           | Non Government | General Medicine      | 2020 | 16                           | 1 Year   |
| 2  | Double-blind, vehicle-controlled, randomised,<br>multi-centre study to evaluate the efficacy and<br>safety of LH-8 cutaneous solution in children<br>and adolescents withmoderate to severe scalp<br>Alopecia Areata.   | Dr. Ashihs Deshmukh  | CliniExperts Research<br>Services Private Limited | Non Government | Dermatology           | 2020 | 2                            | 1 Year   |

| 3 | A multicenter, phase II, double blind,<br>randomized, placebo controlled study to<br>evaluate efficacy of recombinant BCG VPM1002<br>in reducing infection incidence and disease<br>severity of SARS-COV-2/COVID-19 among<br>high risk subjects.  | Dr. Tayade Deepak Narayan   | Serum Institute of India<br>(SII) | Non Government | Community<br>Medicine                 | 2020 | 0.5  | 1 year   |
|---|---|---|-----------------------------------|----------------|---------------------------------------|------|------|----------|
|   | A Multicenter, Phase III, Double-Blind,<br>Randomized, Placebo Controlled Study to<br>Evaluate the Efficacy of Recombinant BCG<br>VPM1002 in Reducing Infection Incidence and<br>Disease Severity of SARS-COV-2/COVID-19<br>Among High-Risk Subject under.( <b>Amended</b> )                                  | Dr. Tayade Deepak Narayan   | DIGNOSEARCH                       | Non Government | Genral Medicine                       | 2020 | 26.4 | 1 Year's |
|   | Promesa DES - 1: A Prospective, Multicenter,<br>Single arm, Open label Study to Evaluate<br>Safety and Performance of Promesa™ DES<br>Sirolimus Eluting Self-Expandable Nitinol<br>Peripheral Stent System for Treating Superficial<br>Femoral Artery (SFA) and Iliac Artery Lesions.                         | Dr. Shivaji Pole  | Meril Life Science Pvt.<br>Ltd.   | Non Government | Radiology                             | 2020 | 1.5  | 1 Year's |
| 5 | Open label, Randomized, Active-controlled,<br>Multi-centric phase II/III Study in Indian<br>Toddlers and Infants to Assess the<br>Immunogenicity and Safety of SIIPL<br>HEXASIILTM (DTwP-HepB-IPV-Hib) Vaccine<br>in Comparison to SIIPL Pentavac (DTwP-HepB-<br>Hib) + Poliovac (IPV) vaccines Administered. | Dr. Deepak Tyade  | HEXA                              | Non Government | Pediatric                             | 2020 | 1.9  | 1 Year's |
| 6 | multicentric phase -II/III study in Indian<br>toddlers and infants to assess the<br>immunogenecity and safety of SIIPL HEXASIIL   | Dr. Tayade Deepak , Dr.<br>Joshi Bhavna Dr.<br>Engade Madhuri Dr.<br>Joshi Pradnya, Dr.<br>Nela Madhura | Serum Institute of India<br>(SII) | Non Government | Community<br>Medicine &<br>Pediatrics | 2020 | 2.5  | 3 year   |
|   | A prospective, multi-centre, open lable, phase<br>IV study to evaluate safety and efficacy profile<br>of Infimab TM in patients with moderate to<br>severe plaque psoriasis.  | Dr Ashish Deshmukh  | Reliance                          | Non Government | Skin & VD                             | 2020 | 7.3  | 1 year   |

| 8  | A prospective, randomized, two-arm, active-<br>controlled, parallel, Multicentre, non-inferiority,<br>phase III clinical trial to assess the efficacy and<br>safety of Amorolfine lotion 0.25% W/v as<br>compared to Amorolfine cream 0.25% w/w in<br>patients with superficial fungal infection of the<br>skin.  | Dr Ashish Deshmukh  | Zydus                       | Non Government | Skin & VD        | 2020 | 2.92 | 2 year |
|----|---|---------------------|-----------------------------|----------------|------------------|------|------|--------|
| 9  | A prospective, Double-Blind, Randomized,<br>Parallel- Group, Vehicle-controlled, Multicentre<br>study to Evaluate the Safety and Bioequivalence<br>of Mupirocin Cream USP 2 % (Supplied<br>By:Glasshouse Pharmaceutical Limited Canada)<br>using Mupirocin Cream USP 2% (Manufactured<br>by: Glenmark Pharmaceuticals Ins; USA) as a<br>Reference product, in subject, with Secondarily<br>Infected Traumatic Skin Lesions. |                     | JSS                         | Non Government | Skin & VD        | 2020 | 0.9  | 1 Year |
| 10 | A Phase III, Multicentric, Prospective,<br>Randomized, Parallel Study to Evaluate the<br>Efficacy and Safety of Molnupiravir in Adult<br>Indian Patients with Mild COVID-19.  | Dr. Umar Quadri     | Hetero Labs Limited         | Non Government | General Medicine | 2021 | 3    | 1 Year |
| 11 | A Phase-III, Multicenter, Prospective, Double<br>Blind, Randomized, Parallel Clinical Study<br>Evaluating The Efficacy, Safety And<br>Tolerability Of Hetero-Tocilizumab In Cytokine<br>Storm Of Severe Coronavirus Disease (Covid-<br>19) Pneumonia (TOCICOVID Study).   | Dr. Anand Nikalje   | Hetero Biopharma<br>Limited | Non Government | General Medicine | 2021 | 23   | 1 Year |
| 12 | A Phase 3, Observer blind, Randomized, Active-<br>Controlled Trial Evaluating the Immunologic<br>Noninferiority, Safety and Tolerability of a 15-<br>valent Pneumococial Conjugate Vaccine<br>Compared to a 13-valent Pneumococcal<br>Conjugate Vaccine in Healthy Infants Given<br>with Routine Pediatric Vaccinations.  | Dr. Mohammed Haseeb | Tergene Biotech Pvt. Ltd    | Non Government | Pediatric        | 2021 | 7    | 1 Year |

| 13 | A multicentre, prospective, open label,<br>randomized comparative clinical trial to<br>evaluate safety and efficacy of Reliance Life<br>Sciences'Bevacizumab (R-TPR-023) plus<br>standard of care and standard of care alone in<br>COVID 19 Acute Respiratory Distress<br>Syndrome (ARDS) Patients with non-invasive<br>ventilation.  | Dr. Anand Nikalje                         | Reliance life Sciences<br>Pvt. Ltd              | Non Government | Genral Medicine | 2021 | 11    | 1 Year   |
|----|---|---|---|----------------|-----------------|------|-------|----------|
|    | A prospective randopmized double blind<br>multicentric two arms phase 3 clinical trial to<br>assess the efficacy and safety of<br>levobupivacaine in dextrose injection<br>Manufactured by Neon Laboratories LTD.<br>Versus that of Bupivacaine heavy (Bupivacaine<br>hydrochloride in dextrose injection<br>Manufactured by Themis Medicare LTD. for<br>spinal Anesthesia in adult patients. | Dr. V. P. Kelkar(PI)<br>Dr. Dhanjay Bhale | CRO- Biosphere<br>Sponsor- Neon<br>Laboratories | Non Government | Anaesthesia     | 2021 | 1.32  | On going |
|    | A multicentre, prospective, open label,<br>randomized comparative clinical trial to<br>evaluate safety and efficacy of Reliance Life<br>Sciences' Bevacizumab (R-TPR-023) plus<br>standard of care and standard of care alone in<br>COVID 19 Acute Respiratory Distress<br>Syndrome (ARDS) Patients with non-invasive<br>ventilation.   | Dr. Anand Nikalje                         | Reliance Life Sciences'                         | Non Government | Genral Medicine | 2021 | 11.15 | 1 Year's |
| 16 | 1   | Dr. Kelkar Vasanti<br>Prabhakar           | Biosphere Clinical<br>Research Pvt. Ltd         | Non Government | Anesthesia      | 2021 | 0.5   | 1 Year   |

|   |  |                     | 2021-2022         |                |              |      |                  |         |
|---|--|---------------------|-------------------|----------------|--------------|------|------------------|---------|
| 1 | Randomized double blinded placebo controlled<br>parralal group multicenter phase II/III adaptive<br>clinical trial to assess the safety and<br>immunogenicity of gam-COVID-vac Combined<br>Vector Vaccine for SARS-Cov-2 Infection in<br>Indian Healthy Subjects   |                     | JSS (Grapecity)   | Non-Government | Medicine     | 2021 | 496059           | 02 Year |
| 2 | A prospective, randomized, two-arm, active-<br>controlled, parallel, multicentre, non-inferiority,<br>phase III clinical trial to assess the efficacy and<br>safety of Amorolfine lotion O.25% w/v as<br>compared to Amorolfine cream 0.25% w/w in<br>patients with superficial fungal infection of the<br>skin.   | Dr. Ashish Deshmukh | Zydus (Grapecity) | Non-Government | Dermatology  | 2021 | 140918<br>59015  | 01 Year |
| 3 | A randomized double blind placebo controlled<br>study evaluating the efficacy & safety of<br>otilimab IV in patient with severe pulmonary<br>COVID 19 related disease.   | Dr. Umar Quedri     | PPD( Doclin C.)   | Non-Government | Medicine     | 2021 | 20000<br>1367500 | 02 Year |
| 4 | A streamlined, multicenter, randomized,<br>parallel group, double blind placebo controlled,<br>superiority trial to evaluate the effect of<br>EMPAgliflozin on hospitalization for heart<br>failure and mortality in patients with acute<br>myocardial infarction.   | Dr. Prashant Udgire | Covance           | Non-Government | Pharmacology | 2021 | 55500            | 01 year |
| 5 | A Prospective, Double-blind, Randomized,<br>Parallel-Group, Vehicle-Controlled, Multicenter<br>Study to Evaluate the Safety and<br>Bioequivalence of Mupirocin Cream USP<br>2% (Supplied by: Glasshouse Pharmaceuticals<br>Limited Canada) using MupirocinCream USP<br>2% (Manufactured by: Glenmark<br>Pharmaceuticals Inc; USA) as a Reference<br>Product, in Subjects with Secondarily Infected<br>Traumatic Skin Lesions | Dr. Ashish Deshmukh | JSS (Grapecity)   | Non Government | Dermatology  | 2021 | 84647            | 2 year  |

| 6  | A multicentre, single arm, phase IV clinical trial<br>to evaluate thesafety and efficacy of Itolizumab<br>for the treatment of cytokinerelease syndrome<br>(CRS) in moderate to severe acute respiratory<br>distress syndrome (ARDS) patients due to<br>COVID 19.   | Dr. Anand Nikaje    | IQVIA (Grapecity)                            | Non Government | Medicine    | 2021 | 153356 | 01 year |
|----|---|---------------------|--|----------------|-------------|------|--------|---------|
| 7  | A Prospective, Multi-centre, Phase IV Study to<br>Assess the Safety, Efficacy and Immunogenicity<br>of BIOSULIN® 30:70 (Insulin Injection, Biphasic<br>Isophane 100 IU/ml of M.J.Biopharm Private<br>Limited) in Treatment of Patients Diagnosed<br>with Type 2 Diabetes Mellitus   | Dr. Umar Quadri     | Abionene                                     | Non-Government | Medicine    | 2021 | 27625  | 01 Year |
| 8  | A randomized double blind placebo controlled<br>study evaluating the efficacy & safety of<br>otilimab IV in patient with severe pulmonary<br>COVID 19 related disease.  | Dr. Umar Quadri     | PPD Pharma<br>(Docclin C.)                   | Non-Government | Medicine    | 2021 | 328557 | 02 year |
| 9  | Safety and Efficacy of Lipiodol® Ultra Fluid in<br>Association with Surgical Glues during<br>Vascular Embolization, a phase IV study  | Dr. Shivaji Pole    | Siro clinpharm (Ardent<br>Clinical Research) | Non Government | Radiology   | 2021 | 94041  | 01 Year |
| 10 | A Prospective, Double-blind, Randomized,<br>Parallel-Group, Vehicle-Controlled, Multicenter<br>Study to Evaluate the Safety and<br>Bioequivalence of Mupirocin Cream<br>USP 2% (Supplied by: Glasshouse<br>Pharmaceuticals Limited Canada) using<br>Mupirocin Cream USP 2% (Manufactured by:<br>Glenmark Pharmaceuticals Inc; USA) as a<br>Reference Product, in Subjects with Secondarily<br>Infected Traumatic Skin Lesions | Dr. Ashish Deshmukh | JSS (Grapecity)                              | Non Government | Dermatology | 2021 | 7389   | 2 Year  |
| 11 | A prospective, randomized, two-arm, active-<br>controlled, parallel, multicentre, non-inferiority,<br>phase lll clinical trial to assess the efficacy and<br>safety of Amorolfine lotion O.2S% w/v as   | Dr. Ashish Deshmukh | Zydus (Grapecity)                            | Non Government | Dermatology | 2021 | 10080  | 2 Year  |

| 12 | A multicentre, phase III, double-blind,<br>randomized, placebo controlled study to<br>evaluate the efficacy of recombinant BCG<br>VPM1002 in reducing infection incidence and<br>disease severity of sars-cov-2/covid-19 among<br>high-risk subjects   | Dr. Deepak Tayade   | Serum (Dignosearch)  | Non Government | Community<br>Medicine | 2021 | 581005.5<br>13500<br>3500 | 01 year |
|----|--|---------------------|----------------------|----------------|-----------------------|------|---------------------------|---------|
| 13 | A prospective, randomized, two-arm, active-<br>controlled, parallel, multicentre, non-inferiority,<br>phase III clinical trial to assess the efficacy and<br>safety of Amorolfine lotion O.2S% w/v as<br>compared to Amorolfine cream 0.25% w/w in<br>patients with superficial fungal infection of the<br>skin.   | Dr. Ashish Deshmukh | Zydus (Grapecity)    | Non Government | Dermatology           | 2021 | 128250                    | 02 year |
| 14 | Randomized, Double-Blind, Placebo-<br>Controlled, Parallel-Group, Multi-Centre Phase<br>II/III Adaptive Clinical Trial to Assess the<br>Safety and Immunogenicity of Gam-COVID-<br>Vac Combined Vector Vaccine for SARS-Cov-2<br>Infection in Indian Healthy Subjects  | Dr. Anand Nikaje    | JSS (Grapecity)      | Non Government | Medicine              | 2021 | 630180                    | 02 year |
| 15 | A Phase III, Multicentre, Randomized, Double<br>Blind, Parallel group, Comparative Clinical<br>Study to evaluate Efficacy and Safety of<br>RopivacaineHydrochloride 0.75% (7.5mg/mL)<br>in Dextrose 8% (80mg/mL) injection compared<br>to Bupivacaine Hydrochloride 0.5% (5mg/mL)<br>in Dextrose 8% (80mg/mL) injection for<br>subjects undergoing lower limb orthopaedic<br>surgeries under spinal anaesthesia. | Dr. Vasanti Kalkar  | Bioshpere            | Non Government | Anaesthesia           | 2021 | 28800                     | 2 year  |
| 16 | A prospective, multi-centre, open label, phase<br>IV studyto evaluate safety and efficacy profile<br>of InfimabTM inpatients with moderate to<br>severe plaque psoriasis   | Dr. Ashish Deshmukh | Reliance (Grapecity) | Non Government | Dermatology           | 2021 | 140036                    | 01 year |

| 17 | A multicentre, single arm, phase IV clinical trial<br>to evaluate the safety and efficacy of Itolizumab<br>for the treatment of cytokinerelease syndrome<br>(CRS) in moderate to severe acute respiratory<br>distress syndrome (ARDS) patients due to<br>COVID 19  | Dr. Anand Nikaje   | Biocon (Grapecity) | Non Government | Department of<br>Medicine | 2021 | 146795 | 01 year |
|----|--|--------------------|--------------------|----------------|---------------------------|------|--------|---------|
| 18 | Randomized double blinded placebo controlled<br>parralal group multicenter phase II/III adaptive<br>clinical trial to assess the safety and<br>immunogenicity of gam-COVID-vac Combined<br>Vector Vaccine for SARS-Cov-2 Infection in<br>Indian Healthy Subjects   |                    | JSS (Grapecity)    | Non Government | Department of<br>Medicine | 2021 | 94770  | 01 year |
| 19 | A multicentre, phase iii, double-blind,<br>randomized, placebo controlled study to<br>evaluate the efficacy of recombinant BCG<br>VPM1002 in reducing infection incidence and<br>disease severity of Sars-Cov-2/Covid-19<br>Among High-Risk Subjects   | Dr. Deepak Tayade  | Serum              | Non Government | Community<br>Medicine     | 2021 | 500    | 01 year |
| 20 | A Multicenter, Prospective, Randomized,<br>Double-Blind, Placebo-Controlled study to<br>evaluate the efficacy and safety of Ketorolac 30<br>mg sublingual tablet in the acute treatment of<br>migraine headache.   | Dr.Manjiri Naik    | Syneos             | Non Government | Medicine                  | 2021 | 83333  | 01 year |
| 21 | RESPIRE - A Randomized, Double-Blind,<br>Placebo-Controlled, Multi-Centre Clinical Trial<br>to Evaluate the Safety and Efficacy of ATR-002<br>in Adult Hospitalized Patients with COVID-19   | Dr, Anand Nikalje  | Clinexel           | Non Government | Medicine                  | 2021 | 83250  | 01 year |
| 22 | A prospective, randomized, parallel,<br>multicentric, phase III clinical trial to assess the<br>efficacy and safety of Molnupiravir 800<br>mgcapsules and standard of care (soc)<br>compared to standard of care (soc) only in<br>patients with polymerase chain reaction (RT-<br>PCR)confirmed Mild Covid-19 infection. | Dr, Anand Nikalje  | Clinenfinity       | Non Government | Medicine                  | 2021 | 104643 | 01 year |
| 23 | A Phase IV, Open-Label, Multi-center Study to<br>Evaluate the Safety of Apixaban in Indian<br>Subjects Undergoing Elective Total Knee<br>Replacement or Total Hip Replacement Surgery  | Dr. Girish Gadekar | PPD Pharma         | Non Government | Orthopedics               | 2021 | 75006  | 01 year |

|    | A prospective, multi-centre, open label, phase<br>IV studyto evaluate safety and efficacy profile<br>of InfimabTM inpatients with moderate to<br>severe plaque psoriasis   | Dr. Ashish Deshmukh | Reliance (Grapecity)         | Non Government | Dermatology           | 2021 | 376866  | 2 year  |
|----|--|---------------------|------------------------------|----------------|-----------------------|------|---------|---------|
| 25 | A multicentre, phase III, double-blind,<br>randomized, placebo controlled study to<br>evaluate the efficacy of recombinant BCG<br>VPM1002 in reducing infection incidence and<br>disease severity of sars-cov-2/covid-19 among<br>high-risk subjects                   | Dr. Deepak Tayade   | Serum (Diagnosearch)         | Non Government | Community<br>Medicine | 2021 | 50000   | 2 year  |
| 26 | A multicentre, phase III, double-blind,<br>randomized, placebo controlled study to<br>evaluate the efficacy of recombinant BCG<br>VPM1002 in reducing infection incidence and<br>disease severity of sars-cov-2/covid-19 among<br>high-risk subjects                   | Dr. Deepak Tayade   | Serum (Diagnosearch)         | Non Government | Community<br>Medicine |      | 1118230 | 01 year |
|    | A multicentre, single arm, phase IV clinical trial<br>to evaluate thesafety and efficacy of Itolizumab<br>for the treatment of cytokine release syndrome<br>(CRS) in moderate to severe acute<br>respiratorydistress syndrome (ARDS) patients<br>due to COVID 19.      | Dr. Anand Nikaje    | Biocon (Grapecity)           | Non Government | Medicine              | 2021 | 30303   | 2 Year  |
| 28 | A prospective, multi-centre, open label, phase<br>IV studyto evaluate safety and efficacy profile<br>of InfimabTM inpatients with moderate to<br>severe plaque psoriasis.  | Dr. Ashish Deshmukh | Reliance (Grapecity)         | Non Government | Dermatology           | 2021 | 77501   | 2 Year  |
| 29 | A Phase-III, Multicenter, Prospective, Double<br>Blind, Randomized, Parallel Clinical Study<br>Evaluating The Efficacy, Safety And<br>Tolerability Of Hetero-Tocilizumab In Cytokine<br>Storm Of Severe Coronavirus Disease (Covid-<br>19) Pneumonia (TOCICOVID Study) | Dr. Anand Nikaje    | HETRO (Grapecity)            | Non Government | Medicine              | 2021 | 1163565 | 2 Year  |
| 30 | Safety and Efficacy of Lipiodol® Ultra Fluid in<br>Association with Surgical Glues during<br>Vascular Embolization, a phase IV study   | Dr. Shivaji Pole    | Siro clinpharm<br>(Reliance) | Non Government | Radiology             | 2021 | 121500  | 2 Year  |

|    | Randomized, Double-Blind, Placebo-<br>Controlled, Parallel-Group, Multi-Centre Phase<br>II/III Adaptive Clinical Trial to Assess the<br>Safety and Immunogenicity of Gam-COVID-<br>Vac Combined Vector Vaccine for SARS-Cov-2<br>Infection in Indian Healthy Subjects | Dr. Anand Nikaje                           | JSS (PPD Pharma)                                | Non Government | Medicine              | 2021 | 25002           | 2 Year           |
|----|---|--|---|----------------|-----------------------|------|-----------------|------------------|
| 32 | Safety and Efficacy of Lipiodol® Ultra Fluid in<br>Association with Surgical Glues during<br>Vascular Embolization, a phase IV study  | Dr. Shivaji Pole                           | Siro clinpharm<br>(Ardent clinical<br>Research) | Non Government | Radiology             | 2021 | 78106           | 2 Year           |
| 33 | A phase two study to evaluate the testicular<br>safety of a nw oral drugs in adult male patient<br>with active alcerative collitties.   | Dr.Ashok Mohite                            | Vedic Life sciences                             | Non Government | Surgery               | 2021 | 74250           | 1 year           |
|    | Randomized, Double-Blind, Placebo-<br>Controlled, Parallel-Group, Multi-Centre Phase<br>II/III Adaptive Clinical Trial to Assess the<br>Safety and Immunogenicity of Gam-COVID-<br>Vac Combined Vector Vaccine for SARS-Cov-2<br>Infection in Indian Healthy Subjects | Dr. Anand Nikaje                           | JSS (Grapecity)                                 | Non Government | Medicine              | 2021 | 82485           | 2 Year           |
|    | Randomized, Double-Blind, Placebo-<br>Controlled, Parallel-Group, Multi-Centre Phase<br>II/III Adaptive Clinical Trial to Assess the<br>Safety and Immunogenicity of Gam-COVID-<br>Vac Combined Vector Vaccine for SARS-Cov-2<br>Infection in Indian Healthy Subjects | Dr. Anand Nikaje                           | JSS   | Non Government | Medicine              | 2021 | 26730           | 3 Year           |
| 36 | A multicentre, phase III, double-blind,<br>randomized, placebocontrolled study to<br>evaluate the efficacy of recombinant BCG<br>VPM1002 in reducing infection incidence and<br>disease severity of Sars-cov-2/covid-19 among<br>high-risk subjects                   | Dr. Deepak Tayade                          | Serum   | Non Government | Community<br>Medicine | 2021 | 12000           | 1 year           |
| 37 | A Randomized. Active-Controlled, Double-<br>Masked, Parallel-Group, Phase 3 Study to<br>Compare Efficacy and Safety of CT-P42 in<br>comparison with Eylea in Patients with Diabetic<br>Macular Edema  | Dr. Deepak Bhosle Dr.<br>Swapnil Dongaokar | Syneos  | Non Government | Pharmacology          | 2021 | 27778           | 1 year           |
| 38 | A Randomized, Double-blind, Parallel Arm,<br>Multicenter Study to Evaluate the Efficacy and<br>Safety of Nitric Oxide Nasal Spray Combined<br>with Standard Supportive Care in Adult<br>non-hospitalized Patients With COVID- 19.                                     | Dr. Anand Nikaje                           | Gelmark (Grapecity)                             | Non Government | Medicine              | 2021 | 550602<br>55890 | 1 year<br>1 year |

| A Phase III Randomized, Double Blind, Parallel<br>Group, Placebo Controlled, Multi-centre,<br>Multinational Study to Evaluate Efficacy and<br>Safety of TRC150094 as an Add On to Standard<br>of Care in Improving Cardiovascular Risk in<br>Subjects with Diabetes, Dyslipidemia and<br>Hypertension. | 1 | IQVIA      | Non Government | Pharmacology | 2021 | 100000.8 | 1 year |
|--|---|------------|----------------|--------------|------|----------|--------|
| A Phase IV, Open-Label, Multi-center Study to<br>Evaluate the Safety of Apixaban in Indian<br>Subjects Undergoing Elective Total Knee<br>Replacement or Total Hip Replacement Surgery  |   | PPD Pharma | Non Government | Orthopedics  | 2021 | 50001    | 1 year |

|   |  | 2022-2023   |  |                |  |      |          |        |
|---|--|---|--|----------------|--|------|----------|--------|
| 1 | A Multidisciplinary Educational training course for physical therapists on<br>managing common musculoskeletal pain conditions in Maharashtra, India  | PI: Dr Rinkle Malani<br>CO PI: Dr Prashant<br>Mukkannavar<br>Dr Tajuddin Chitapure<br>Dr Vaibhavi Walimbe | International<br>Association<br>for the study<br>of pain |                | MGM School of<br>Physiotherapy<br>Aurangabad | 2022 | 6.74206  | 1 Year |
| 2 | A phase two study to evaluate the testicular safety of a nw oral drugs in<br>adult male patient with active alcerative collitties.   | Dr. Pravin Suryawanshi  | Vedic Life<br>sciences                                   | Non Government | Surgery                                      | 2022 | 0.03459  | 1 Year |
| 3 | A Randomized, Double-Blind, Double-Dummy, Multiple-Dose,<br>Multicenter, Three-Arm, Parallel Study to Compare the Efficacy and<br>Safety of Pregabalin ER Tablet of Alvogen Malta (Out-Licensing) Ltd. to<br>Placebo and Lyrica® (Pregabalin) Hard Capsule of Pfizer in Subjects with<br>Diabetic Peripheral Neuropathy.   | Dr.Deepak Bhosle  | Cliantha   | Non-Government | Pharmacology                                 | 2022 | 0.29174  | 1 year |
| 4 | A Randomized, Single Blind, Placebo Controlled, Phase 2 Clinical Trial to<br>Evaluate the Efficacy and Safety of Ursodiol Injection 625mg/25ml<br>(25mg/ml) in Adults with moderate COVID 19 patients at a dose of 1750<br>mg/day via intravenous infusion   | Dr. Rohan Gundre  | Abiogenesis  | Non-Government |  | 2022 | 9.01848  | 1 year |
| 5 | Study Title: A Randomized, Open-label, Prospective, Comparative, Multi-<br>centre, Parallel group, Active controlled, Phase III Study to Evaluate the<br>Efficacy and Safety of the Ropivacaine Readyfusor 2mglml versus<br>Ropivacaine Balloon Pump Infusor 2mglml as a continuous surgical site<br>infusion for the Treatment of Post-Surgicat Pain in lower abdominal<br>laparotomy | Dr.Vasanti kelkar   | JSS medical  | Non-Government | Anaesthesia                                  | 2022 | 0.225    | 1 year |
| 6 | A Phase 3, Multicenter, Randomized, Double-blind, Placebo-controlled<br>Trial to<br>Evaluate the Efficacy and Safety of Sibeprenlimab Administered<br>Subcutaneously in Subjects with<br>Immunoglobulin A Nephropathy- Study Number 417 201 00007  | Dr.Sudhir Kulkarni  | George   | Non-Government | Nephrology                                   | 2022 | 1.500012 | 1 year |
| 7 | A randomized double blind placebo controlled study evaluating the efficacy & safety of otilimab IV in patient with severe pulmonary COVID 19 related disease.  | Dr.Syed Umar Qudri  | Doclin   | Non-Government | medicine                                     | 2022 | 0.97308  | 1 year |
| 8 | A multicentre, phase III, double-blind, randomized, placebo controlled<br>study to evaluate the efficacy of recombinant BCG VPM1002 in reducing<br>infection incidence and disease severity of sars-cov-2/covid-19 among<br>high-risk subjects   | Dr.Deepak tayade  | Dignosearch  | Non-Government |  | 2022 | 4.81258  | 1 year |
| 9 | A prospective, randomized, parallel, multicentric, phase III clinical trial to<br>assess the efficacy and safety of Molnupiravir 800 mgcapsules and<br>standard of care (soc) compared to standard of care (soc) only in patients<br>with polymerase chain reaction (RT-PCR)confirmed Mild Covid-19<br>infection.  | Dr.Syed Umar Qudri  | Cliniinfinity  | Non-Government | Medicine                                     | 2022 | 4.13739  | 1 year |

| 10 | KETO-22-002/" A Randomized, Double Blind, Placebo Controlled, Parallel<br>Design, Multi-Center, Bioequivalence Study with Clinical<br>Endpoint of Ketoconazole Shampoo 2% (Test Product)<br>Compared with Ketoconazole Shampoo 2% (Reference  | Dr.Ashish Deshmukh      | Scitus pharma                      | Non-Government | skin        | 2022 | 0.85    | 1 year |
|----|---|-------------------------|------------------------------------|----------------|-------------|------|---------|--------|
|    | Product) In Human Adult Male and Female (non-pregnant)<br>Subjects with Tinea Versicolor."  |                         |                                    |                |             |      |         |        |
| 11 | A Prospective, Multi-centre, Active Post-Marketing Surveillance Study of<br>Liposomal Amphotericin B (Amphonex® - Manufactured by Bharat<br>serums and Vaccines ltd) in Patients with Invasive Fungal infection who<br>are refractory to or intolerant of conventional Amphotericin B therapy in<br>Real World Clinical Practice (AMBeR).   | Dr.Anand Nikalje        | synegene                           | Non-Government | Medicine    | 2022 | 0.0002  | 1 year |
| 12 | A Phase 3, Observer blind, Randomized, Active-Controlled Trial<br>Evaluating the Immunologic Non-inferiority, Safety and Tolerability of a<br>15-valent Pneumococcal Conjugate Vaccine Compared to a 13-valent<br>Pneumococcal Conjugate Vaccine in Healthy Infants Given with Routine<br>Pediatric Vaccinations.   | Dr.haseeb               | Grapecity                          | Non-Government | Pediatrics  | 2022 | 1.02141 | 1 year |
| 13 | A Phase 3, Observer blind, Randomized, Active-Controlled Trial<br>Evaluating the Immunologic Non-inferiority, Safety and Tolerability of a<br>15-valent Pneumococcal Conjugate Vaccine Compared to a 13-valent<br>Pneumococcal Conjugate Vaccine in Healthy Infants Given with Routine<br>Pediatric Vaccinations.   | Dr.haseeb               | Grapecity                          | Non-Government | Pulmonary   | 2022 | 0.5499  | 1 year |
| 14 | A Randomized, Double-blind, Parallel, Placebo Controlled, Three Arm,<br>Multicentric Study to Evaluate the Efficacy and Safety of Ketoconazole 2%<br>Cream of Cadila Healthcare Limited with Ketoconazole 2% Cream of Teva<br>Pharmaceuticals USA in Subjects with Tinea Pedis.   | Dr.Ashish Deshmukh      | Cadila<br>Healthcare<br>(Cliantha) | Non-Government | Dermatology | 2022 | 1.76715 | 1 year |
| 15 | CPB-007-2021<br>CPB-009-2021<br>CPB-010-2021<br>CPB-005-2021  | Dr. Sai Kumar Gowalikar | Wockhardt R<br>& D                 | Non-Government |             | 2022 | 1.35    | 1 year |
| 16 | A Prospective, Multi-centre, Active Post-Marketing Surveillance Study of<br>Liposomal Amphotericin B (Amphonex® - Manufactured by Bharat<br>serums and Vaccines ltd) in Patients with Invasive Fungal infection who<br>are refractory to or intolerant of conventional Amphotericin B therapy in<br>Real World Clinical Practice (AMBeR).   | Dr.Anand Nikalje        | Synegene                           | Non-Government |             | 2022 | 0.75    | 1 year |
| 17 | A Multicenter, Randomized, Parallel-Group, 6-Week Treatment Clinical<br>Study to Assess Bioequivalence of Budesonide 80 µg and Formoterol<br>Fumarate Dihydrate 4.5 µg Inhalation Product (Cipla Ltd.) in comparison<br>with the Reference Product, Symbicort® (Budesonide/Formoterol<br>Fumarate Dihydrate, 80/4.5 µg per Actuation) Inhalation Aerosol<br>(AstraZeneca, USA), in Adult Asthma Patients. | Dr.hafiz Deshmukh       | Parexel                            | Non-Government | Pulmonory   | 2022 | 0.75    | 1 year |
| 18 | A Randomized Double-blind, Four-Arm Active and Placebo-controlled<br>DoseFinding Trial to Evaluate the Efficacy, Tolerability, Safety and Dose<br>Response of LYT-100 in Patients with Idiopathic Pulmonary Fibrosis (IPF).   | Dr.hafiz Deshmukh       | Novotech                           | Non-Government | pulmonary   | 2022 | 0.75    | 1 year |

| 19 | A Prospective, Multi-centre, Active Post-Marketing Surveillance Study of<br>Liposomal Amphotericin B (Amphonex® - Manufactured by Bharat<br>serums and Vaccines ltd) in Patients with Invasive Fungal infection who<br>are refractory to or intolerant of conventional Amphotericin B therapy in<br>Real World Clinical Practice (AMBeR).   | Dr.Anand Nikalje   | Bharat Serume<br>Vaccine(<br>Syngene) | Non-Government | Medicine     | 2022 | 0.3     | 1 year |
|----|---|--------------------|---------------------------------------|----------------|--------------|------|---------|--------|
| 20 | An open label, randomized, active controlled, multi-centric Phase II/III,<br>study in Indian toddlers and infants to assess the immunogenicity and<br>safety of SIIPL HEXASIIL <sup>TM</sup> (DTwP-HepB-IPV-Hib) Vaccine in<br>comparison with SIIPL PENTAVAC (DTwP-HepB-Hib) + Poliovac (IPV)<br>vaccines administered as separate injections.   | Dr.Deepak tayade   | Serum<br>Institute (<br>Diagosearch)  | Non-Government | РСМ          | 2022 | 0.76668 | 1 year |
| 21 | Randomized, Multicenter, Single-dose, Two-treatment, Single-period,<br>Parallel, Bioequivalence Study of Mylan's Paliperidone Prolonged release<br>Suspension for Injection, 175 mg and Janssen-Cilag's Trevicta® prolonged<br>release suspension for injection, 175 mg in Subjects withSchizophrenia   | Dr.manik Bhise     | Siro clinpharm                        | Non-Government | Psychiatry   | 2022 | 0.85    | 1 year |
| 22 | International multicenter comparative prospective single-blind study of<br>efficacy, safety, pharmacodynamics, pharmacokinetics and<br>immunogenicity of the drug product Tenecteplase, lyophilized for<br>solution for intravenous injection 50 mg (Pharmasyntez-Nord JSC, Russia)<br>and the drug product Metalyse®, lyophilized for solution for intravenous<br>injection 50 mg (Boehringer Ingelheim International GmbH, Germany) in<br>patients with ST-segment elevation myocardial infarction (STEMI). | Dr.Prashant udgire | JSS medical<br>/Pharmacenzt           | Non-Government | Cardiology   | 2022 | 0.765   | 1 year |
| 23 | A Randomized, Open-Label, Parallel-Group Study To Evaluate The Safety<br>And Efficacy of Abrocitinib 100 Mg And 200 Mg Tablets In Participants<br>Aged 12 Years And Older With Moderate To Severe Atopic Dermatitis In<br>India."   | Dr.Ashish Deshmukh | ICON                                  | Non-Government | Dermatology  | 2022 | 0.3     | 1 year |
| 24 | Double-blind, vehicle-controlled, randomised, multi-centre study to<br>evaluate the efficacy and safety of LH-8 cutaneous solution in children<br>and adolescents with moderate to severe scalp Alopecia Areata   | Dr.Ashish Deshmukh | Grapecity                             | Non-Government | Dermatology  | 2022 | 0.85556 | 1 year |
| 25 | A Randomized, Double-blind, Parallel, Placebo Controlled, Three Arm,<br>Multicentric Study to Evaluate the Efficacy and Safety of Ketoconazole 2%<br>Cream of Cadila Healthcare Limited with Ketoconazole 2% Cream of Teva<br>Pharmaceuticals USA in Subjects with Tinea Pedis."  | Dr.Ashish Deshmukh | Cliantha                              | Non-Government |              | 2022 | 0.7956  | 1 year |
| 26 | "A 26 Week, Multicenter, Randomized, Placebo-Controlled, Double-Blind,<br>Parallel Group, Phase 3 Trial with a 26 Week Safety Extension Period<br>Evaluating the Safety and Efficacy of Dapagliflozin 5 and 10 mg, and<br>Saxagliptin 2.5 and 5 mg in Pediatric Patients with Type 2 Diabetes   | Dr.Deepak Bhosle   | Grapecity                             | Non-Government | Pharmacology | 2022 | 1.28143 | 1 year |

| 27 | A Randomized Double-blind, Four-Arm Active and Placebo-controlled<br>DoseFinding Trial to Evaluate the Efficacy, Tolerability, Safety and Dose<br>Response of LYT-100 in Patients with Idiopathic Pulmonary Fibrosis (IPF).   | Dr.hafiz Deshmukh   | Novotech                 | Non-Government | Pulmonary          | 2022 | 0.3     | 2 year |
|----|---|---------------------|--------------------------|----------------|--------------------|------|---------|--------|
| 28 | "A bioequivalence study of Levothyroxine sodium tablets 100 mcg of<br>Wockhardt Limited, India compared with Eltroxin® 100 mcg<br>(Levothyroxine sodium) tablets of Mercury Pharma Group Ltd., UK in<br>healthy, adult, human subjects under fasting condition."  | Sai Kumar Gowalikar | Wockhardt R<br>& D       | Non-Government | Wockhardt R &<br>D | 2022 | 0.3375  | 2 year |
| 29 | A Randomized Double-blind, Four-Arm Active and Placebo-controlled<br>DoseFinding Trial to Evaluate the Efficacy, Tolerability, Safety and Dose<br>Response of LYT-100 in Patients with Idiopathic Pulmonary Fibrosis (IPF).   | Dr.hafiz Deshmukh   | Novotech                 | Non-Government | Pulmonary          | 2022 | 0.3     | 1 year |
| 30 | KETO-22-002/ A Randomized, Double Blind, Placebo Controlled, Parallel<br>Design, Multi-Center, Bioequivalence Study with Clinical Endpoint of<br>Ketoconazole Shampoo 2% (Test Product) Compared with Ketoconazole<br>Shampoo 2% (Reference Product) In Human Adult Male and Female (non-<br>pregnant) Subjects with Tinea Versicolor | Dr.Ashish Deshmukh  | Oxygenclin               | Non-Government | Dermatology        | 2022 | 1.23433 | 1 year |
| 31 | A Randomized, Double-blind, Parallel, Placebo Controlled, Three Arm,<br>Multicentric Study to Evaluate the Efficacy and Safety of Ketoconazole 2%<br>Cream of Cadila Healthcare Limited with Ketoconazole 2% Cream of Teva<br>Pharmaceuticals USA in Subjects with Tinea Pedis.   | Dr.Ashish Deshmukh  | Cliantha                 | Non-Government | Dermatology        | 2022 | 0.0945  | 1 year |
| 32 | A randomized, open-label, parallel-group study to evaluate the safety and efficacy of Abrocitinib 100 mg and 200 mg tablets in participants aged 12 years and older with moderate to severe atopic dermatitis in India  | Dr.Ashish Deshmukh  | Abad HC &<br>Researh LLP | Non-Government | Dermatology        | 2022 | 2.27662 | 2 year |
| 33 | Safety and Efficacy of Transarterial Chemoembolization with Lipiodol® in<br>the treatment of inoperable Hepatocellular Carcinoma (HCC) in Indian<br>Patients: Phase IV Clinical Trial.  | Dr.Shivaji Pole     | Siro                     | Non-Government | Radiology          | 2022 | 0.31068 | 2 year |

|   |   |                           | 2023-2024  |                |                 |      |        |         |
|---|---|---------------------------|--|----------------|-----------------|------|--------|---------|
| 1 | Evaluating the accuracy and efficacy of an<br>indigenously developed speech-to text software<br>(Digisehat) in general radiology reports in a<br>hospital setting   | Dr. Chandramani<br>Pathak | Jatayu<br>Healthcare<br>Technologies<br>(Mumbai) | Non Government | Biotechnology   | 2023 | 125000 | 2 Years |
| 2 | Validation of Biochemical Assays: Extraction of DNA and RNA from Clinical samples   | Dr. Chandramani<br>Pathak | MP Biomedical<br>India Pvt. Ltd.                 | Non Government | Biotechnology   | 2023 | 270000 | 2 Years |
| 3 | Assessment of drug utilization patterns of oral<br>cephalexin-clavulanic acid combination and other<br>antibiotics for the management of surgical site<br>infections  | Dr.Abdul Qayyum<br>Khan   | Meticulous<br>Healthcare                         | Non Government | Surgery         | 2023 | 31590  | 1 Year  |
| 4 | A randomized, open-label, parallel-group study<br>to evaluate the safety and efficacy of abrocitinib<br>100 mg and 200 mg tablets in participants aged 12<br>years and older with moderate to severe atopic<br>dermatitis in India.   | Dr.Ashish Deshmukh        | Aurangabad<br>Healthcare<br>Research (SMO)       | Non Government | Skin and VD     | 2023 | 227662 | 1 Year  |
| 5 | A Randomized, Double-blind, Parallel, Placebo<br>Controlled, Three Arm, Multicentric Study to<br>Evaluate the Efficacy and Safety of Ketoconazole<br>2% Cream of Cadila Healthcare Limited with<br>Ketoconazole 2% Cream of Teva Pharmaceuticals<br>USA in Subjects with Tinea Pedis. | Dr.Ashish Deshmukh        | Cliantha   | Non Government | Skin and VD     | 2023 | 45000  | 1 Year  |
| 6 | A prospective, interventional, open-label, phase<br>1a/1b, single- center,<br>clinical study to evaluate the safety and<br>tolerability of NV-CoV-2 Oral Syrup and Oral<br>Gummy (Chewable gel) for oral administration in  | Dr.Anand Nikalje          | Karveer<br>Meditech                              | Non Government | Medicine        | 2023 | 195122 | 1 Year  |
| 7 | Cipla Ltd, CRD/20, A Multicenter, Randomized,<br>Parallel-Group, 6-Week Treatment Clinical Study<br>to Assess Bioequivalence of Budesonide 80 µg and<br>Formoterol Fumarate Dihydrate 4.5 µg Inhalation<br>Product (Cipla Ltd.) in comparison with the                                | Dr.Hafiz Deshmukh         | Grapecity  | Non Government | Pulmonary Medic | 2023 | 83289  | 1 Year  |
| 8 | Cipla Ltd, CRD/20, A Multicenter, Randomized,<br>Parallel-Group, 6-Week Treatment Clinical Study<br>to Assess Bioequivalence of Budesonide 80 µg and<br>Formoterol Fumarate Dihydrate 4.5 µg Inhalation<br>Product (Cipla Ltd.) in comparison with the                                | Dr.Hafiz Deshmukh         | Grapecity  | Non Government | Pulmonary Medic | 2023 | 24863  | 1 Year  |

| 9  | Cipla Ltd, CRD/20, A Multicenter, Randomized,<br>Parallel-Group, 6-Week Treatment Clinical Study    | Dr.Hafiz Deshmukh     | Grapecity       | Non Government   | Pulmonary Medic   | 2023 | 333666 | 1 Year |
|----|---|-----------------------|-----------------|------------------|-------------------|------|--------|--------|
|    | to Assess Bioequivalence of Budesonide 80 µg and<br>Formoterol Fumarate Dihydrate 4.5 µg Inhalation |                       |                 |                  |                   |      |        |        |
| 10 | Product (Cipla Ltd.) in comparison with the<br>KETO-22-002/"A Randomized, Double Blind,             | Dr.Ashish Deshmukh    | Scitus          | Non Government   | Skin and VD       | 2023 | 20250  | 1 Year |
| 10 | Placebo Controlled, Parallel  | D1.7 ISHISH DCSHIIUKH | Seitus          | Non Government   |                   | 2023 | 20250  | i icai |
|    | Design, Multi-Center, Bioequivalence Study with   |                       |                 |                  |                   |      |        |        |
|    | Clinical  |                       |                 |                  |                   |      |        |        |
|    | Endpoint of Ketoconazole Shampoo 2% (Test   |                       |                 |                  |                   |      |        |        |
| 11 | A Randomized, Double-blind, Parallel, Placebo   | Dr.Ashish Deshmukh    | Cliantha        | Non Government   | Skin and VD       | 2023 | 180000 | 1 Year |
|    | Controlled, Three Arm, Multicentric Study to  |                       |                 |                  |                   |      |        |        |
|    | Evaluate the Efficacy and Safety of Ketoconazole  |                       |                 |                  |                   |      |        |        |
|    | 2% Cream of Cadila Healthcare Limited with  |                       |                 |                  |                   |      |        |        |
|    | Ketoconazole 2% Cream of Teva Pharmaceuticals   |                       |                 |                  |                   |      |        |        |
| 12 | KETO-22-002/"A Randomized, Double Blind,  | Dr.Ashish Deshmukh    | Scitus          | Non Government   | Skin and VD       | 2023 | 24300  | 1 Year |
|    | Placebo Controlled, Parallel  |                       |                 |                  |                   |      |        |        |
|    | Design, Multi-Center, Bioequivalence Study with   |                       |                 |                  |                   |      |        |        |
|    | Clinical  |                       |                 |                  |                   |      |        |        |
|    | Endpoint of Ketoconazole Shampoo 2% (Test   |                       |                 |                  |                   |      | •••••  | 1.3/   |
| 13 | Mozec PEB PTA PK Study: A prospective, single-  | DrShivaji Pole        | Ardent Clinical | Non Government   |                   | 2023 | 389887 | 1 Year |
|    | arm, multi-centre, pharmacokinetic study to   |                       | Research        |                  | Radiologist       |      |        |        |
|    | evaluate human pharmacokinetics of the drug,  |                       |                 |                  |                   |      |        |        |
|    | paclitaxel in MozecTM PEB for PTA   |                       |                 |                  |                   |      |        |        |
| 14 | KETO-22-002/"A Randomized, Double Blind,  | Dr.Ashish Deshmukh    | Oxygen Clinical | Non Government   | Skin and VD       | 2023 | 141405 | 1 Year |
|    | Placebo Controlled, Parallel  |                       | Research SMO    |                  |                   | _0_0 |        |        |
|    | Design, Multi-Center, Bioequivalence Study with   |                       |                 |                  |                   |      |        |        |
|    | Clinical  |                       |                 |                  |                   |      |        |        |
|    | Endpoint of Ketoconazole Shampoo 2% (Test   |                       |                 |                  |                   |      |        |        |
| 15 | A randomized, assessor-blind, placebo   | Dr.Hafiz Deshmukh     | Grapecity       | Non Government   | Pulmonary Medic   | 2023 | 245700 | 1 Year |
|    | controlled,multi  |                       |                 |                  |                   |      |        |        |
|    | center, clinical endpoint bioequivalence study to   |                       |                 |                  |                   |      |        |        |
|    | compare the efficacy and safety to generic  |                       |                 |                  |                   |      |        |        |
|    | fluticasone propionate inhalation aerosol USP 44  | D 11 (1 D 1 11        |                 |                  |                   |      |        |        |
| 16 | A randomized, assessor-blind, placebo   | Dr.Hafiz Deshmukh     | Grapecity       | Non Government   | Pulmonary Medic   | 2023 | 312266 | 1 Year |
|    | controlled,multi  |                       |                 |                  |                   |      |        |        |
|    | center, clinical endpoint bioequivalence study to   |                       |                 |                  |                   |      |        |        |
|    | compare the efficacy and safety to generic  |                       |                 |                  |                   |      |        |        |
| 17 | fluticasone propionate inhalation aerosol USP 44<br>A randomized. Doubu-buno. Four-arm active ano   | Dr.Hafiz Deshmukh     | Aurangabad      | Non Covernment   | Pulmonary Medic   | 2023 | 95000  | 1 Year |
| 1/ | racedo.controued dose.finding trial to evaluate   |                       | Healthcare      | TNOIL GOVERNment | i unitonary weatc | 2023 | 90000  | 1 1001 |
|    | efficacy. Tolerabiuty. Safety and dose response of  |                       | Research (SMO)  |                  |                   |      |        |        |
|    | lyt.ioo in patients with ioiopat»ec pulmonary   |                       | (SiviO)         |                  |                   |      |        |        |
|    | fibrosis  |                       |                 |                  |                   |      |        |        |

| 18 | An Open label, Randomized, Active-controlled,<br>Multi-centric Phase-II/III Study in Indian<br>Toddlers and Infants to Assess the<br>Immunogenicity and Safety of SIIPL HEXASIIL <sup>TM</sup><br>(DTwP-HepB-IPV-Hib) Vaccine in Comparison<br>with SIIPL Pentavac (DTwP-HepB-Hib) +<br>Poliovac (IPV) Vaccines, Administered as<br>Separate Injections  | Dr. Deepak Tayade        | Diagnosearch                    | Non Government | Community<br>Medicine | 2023 | 67500  | 1 Year |
|----|--|--------------------------|---------------------------------|----------------|-----------------------|------|--------|--------|
| 19 | Cipla Ltd, CRD/20, A Multicenter, Randomized,<br>Parallel-Group, 6-Week Treatment Clinical Study<br>to Assess Bioequivalence of Budesonide 80 µg and<br>Formoterol Fumarate Dihydrate 4.5 µg Inhalation<br>Product (Cipla Ltd.) in comparison with the<br>Reference Product, Symbicort®<br>(Budesonide/Formoterol Fumarate Dihydrate,<br>80/4.5 µg per Actuation) Inhalation Aerosol<br>(AstraZeneca, USA), in Adult Asthma Patients | Dr.Hafiz Deshmukh        | Grapecity                       | Non Government | Pulmonary Medic       | 2023 | 189482 | 1 Year |
| 20 | Cipla Ltd, CRD/20, A Multicenter, Randomized,<br>Parallel-Group, 6-Week Treatment Clinical Study<br>to Assess Bioequivalence of Budesonide 80 µg and<br>Formoterol Fumarate Dihydrate 4.5 µg Inhalation<br>Product (Cipla Ltd.) in comparison with the   | Dr.Hafiz Deshmukh        | Grapecity                       | Non Government | Pulmonary Medic       | 2023 | 20477  | 1 Year |
| 21 | Comparative clinical evaluation of Efficacy and<br>Safety of Clotrimazole Vaginal<br>film vs Canesten V6 Vaginal Tablet In The<br>Management Of Symptomatic Vulvovaginal<br>candidiasis in non-pregnant women Open   | Dr.Lakshmi<br>Rachakonda | Grapecity                       | Non Government | OBGY                  | 2023 | 171113 | 1 Year |
| 22 | Prospective, Multi-centre, Active Post-Marketing<br>Surveillance Study of Liposomal Amphotericin B<br>(Amphonex® - Manufactured by Bharat serums<br>and Vaccines ltd) in Patients with Invasive Fungal<br>infection who are refractory to or intolerant of   | Dr.Anand Nikalje         | Grapecity                       | Non Government | Medicine              | 2023 | 81907  | 1 Year |
| 23 | KETO-22-002/"A Randomized, Double Blind,<br>Placebo Controlled, Parallel Design, Multi-Center,<br>Bioequivalence Study with Clinical Endpoint of<br>Ketoconazole Shampoo 2% (Test Product)<br>Compared with Ketoconazole Shampoo 2%  | Dr.Ashish Deshmukh       | Scitus                          | Non Government | Skin and VD           | 2023 | 41580  | 1 Year |
| 24 | KETO-22-002/" A randomized, double blind,<br>placebo controlled, parallel design, multi-centre,<br>bioequivalence study with clinical endpoint of<br>ketoconazole shampoo 2% (test product)<br>compared with ketoconazole shampoo 2%   | Dr.Ashish Deshmukh       | Oxygen Clinical<br>Research SMO | Non Government | Skin and VD           | 2023 | 222430 | 1 Year |

| 25 | Parallel-Group, 6-Week Treatment Clinical Study<br>to Assess Bioequivalence of Budesonide 80 µg and<br>Formoterol Fumarate Dihydrate 4.5 µg Inhalation  | Dr.Hafiz Deshmukh  | Grapecity                                  | Non Government | Pulmonary Medic | 2023 | 174783 | 1 Year |
|----|---|--------------------|--|----------------|-----------------|------|--------|--------|
| 26 | Product (Cipla Ltd.) in comparison with the<br>A randomized, assessor-blind,placebo<br>controlled,multi<br>center,clinical endpoint bioequivalence study to<br>compare the efficacy and safety to generic   | Dr.Hafiz Deshmukh  | Grapecity                                  | Non Government | Pulmonary Medic | 2023 | 73710  | 1 Year |
| 27 | fluticasone propionate inhalation aerosol USP 44<br>Cipla Ltd, CRD/20, A Multicenter, Randomized,<br>Parallel-Group, 6-Week Treatment Clinical Study<br>to Assess Bioequivalence of Budesonide 80 µg and<br>Formoterol Fumarate Dihydrate 4.5 µg Inhalation   | Dr.Hafiz Deshmukh  | Grapecity                                  | Non Government | Pulmonary Medic | 2023 | 574850 | 1 Year |
| 28 | Product (Cipla Ltd.) in comparison with the<br>Cipla Ltd, CRD/20, A Multicenter, Randomized,<br>Parallel-Group, 6-Week Treatment Clinical Study<br>to Assess Bioequivalence of Budesonide 80 µg and<br>Formoterol Fumarate Dihydrate 4.5 µg Inhalation  | Dr.Hafiz Deshmukh  | Grapecity                                  | Non Government | Pulmonary Medic | 2023 | 37664  | 1 Year |
| 29 | Product (Cipla Ltd.) in comparison with the<br>A Phase 3, Multicenter, Randomized, Double-<br>blind, Placebo-controlled Trial to<br>Evaluate the Efficacy and Safety of Sibeprenlimab<br>Administered Subcutaneously in Subjects with   | Dr.Sudhir Kulkarni | George Clinical                            | Non Government | Nephrology      | 2023 | 47266  | 1 Year |
| 30 | Parallel-Group, 6-Week Treatment Clinical Study<br>to Assess Bioequivalence of Budesonide 80 µg and<br>Formoterol Fumarate Dihydrate 4.5 µg Inhalation  | Dr.Hafiz Deshmukh  | Grapecity                                  | Non Government | Pulmonary Medic | 2023 | 48350  | 1 Year |
| 31 | Product (Cipla Ltd.) in comparison with the<br>Cipla Ltd, CRD/20, A Multicenter, Randomized,<br>Parallel-Group, 6-Week Treatment Clinical Study<br>to Assess Bioequivalence of Budesonide 80 µg and<br>Formoterol Fumarate Dihydrate 4.5 µg Inhalation<br>Product (Cipla Ltd.) in comparison with the | Dr.Hafiz Deshmukh  | Grapecity                                  | Non Government | Pulmonary Medic | 2023 | 447103 | 1 Year |
| 32 |   | Dr.Hafiz Deshmukh  | Grapecity                                  | Non Government | Pulmonary Medic | 2023 | 30078  | 1 Year |
|    |   | Dr.Ashish Deshmukh | Aurangabad<br>Healthcare<br>Research (SMO) | Non Government | Skin and VD     | 2023 | 107499 | 1 Year |

| 34 | A randomized double-blind, placebo-controlled,<br>multicenter trial assessing the impact of<br>lipoprotein (a) lowering with TQJ230 on major<br>cardiovascular events in patients with established<br>cardiovascular disease.  | Dr.Prashant Udgire  | Novartis   | Non Government | Cardiology   | 2023 | 16848  | 1 Year   |
|----|--|---|--|----------------|--|------|--------|----------|
| 35 | A randomized double-blind, placebo-controlled,<br>multicenter trial assessing the impact of<br>lipoprotein (a) lowering with TQJ230 on major<br>cardiovascular events in patients with established<br>cardiovascular disease.  | Dr.Prashant Udgire  | Novartis   | Non Government | Cardiology   | 2023 | 90000  | 1 Year   |
| 36 | A randomized, double blind, three-arm, placebo<br>controlled, parallel design, multi-center, clinical<br>end point bioequivalence study of ketoconazole<br>shampoo 2% (test product) compared with<br>ketoconazole shampoo 2% (reference product) in<br>adult subjects with tinea versicolor.  | Dr.Ashuish Deshmukh   | Oxygen Clinical<br>Research SMO                        | Non Government | Skin and VD  | 2023 | 51675  | 1 Year   |
| 37 | Comparative clinical evaluation of efficacy and<br>safety of clotrimazole vaginal film vs canesten v6<br>vaginal tablet in the management of symptomatic<br>vulvovaginal candidiasis in non-pregnant women.<br>- open label, randomized, parallel, prospective,<br>multicentric study.   | Dr.Lakshmi<br>Rachakonda                                      | Grapecity SMO  | Non Government | OBGY   | 2023 | 200363 | 1 Year   |
|    | A randomized, assessor-blind,placebo<br>controlled,multicenter,clinical endpoint<br>bioequivalence study to compare the efficacy and<br>safety to generic fluticasone propionate inhalation<br>aerosol USP 44 mcg (Glenmark pharmaceutical<br>Ltd) to flovent<br>HFA (Fluticasone propionate inhalation<br>aerosol)44 mcg (GSK Group of companies) in<br>treatment of patient with bronchial asthma. | Dr.Hafiz Deshmukh   | Grapecity  | Non Government | Pulmonary Medic  | 2023 | 167310 | 1 Year   |
| 39 | Rotasil@ vaccine impact assessment study in<br>Kerala , Maharashtra and Gurjarat, India  | Dr. Mohad Haseeb  | CMC<br>VELLOR<br>Vellor                                | Non Government | Pediatircs   | 2023 | 288120 | 2 Years  |
| 40 | Influence of expiratory yoga techniques on<br>respiratory function, physical activity, quality of<br>life, and length of hospital stay in patients with<br>pleural disorders: A Randomized Controlled Trial  | Shwelita Mehta Bela<br>Agarwal                                | Society of<br>Indian<br>Physiotherapist                | Non Government | MGM School of<br>Physiotherapy<br>Navi Mumbai              | 2023 | 15000  | 2 Years  |
| 41 | Compare the difference between traditional<br>surveys and drone surveys for identification of<br>vector breeding places at high risk areas in<br>Aurangabad city Maharashtra   | Dr.Shivcharan<br>Kantarao Kendre<br>Dr.<br>Shobha Banci Salve | Indian public<br>Health<br>Association-<br>Maharashtra | Non Government | Department of<br>Community,<br>MGM Medical<br>College, ABD | 2024 | 15000  | 6 months |

| 42 | Assessment of Health Seeking Behaviour and | Dr. Pahune V. G. | Indian public | Non Government | Department of | 2023 | 6000 | 6 months |
|----|--|------------------|---------------|----------------|---------------|------|------|----------|
|    | health Profile among Transgenders in       | Dr. Tayade       | Health        |                | Community,    |      |      |          |
|    | Aurangabad city - a cross sectional study  | D.N              | Association-  |                | MGM Medical   |      |      |          |
|    |  | Dr. Mahajan S.M. | Maharashtra   |                | College, ABD  |      |      |          |
|    |  | Dr Joshi         |               |                |               |      |      |          |
|    |  | B.P              |               |                |               |      |      |          |