

FTS No: 21962
Date: 25.04.2018

F. No 12-24/12-DC
Government of India
Directorate General of Health Services
Central Drugs Standard Control Organisation
(New Drugs Division)

Tele No.011-23236965
Fax.No.011-23236973

FDA Bhawan, Kotla Road
New Delhi
Dated 05.05.2019

To
M/s Urshapharm India Private Limited,
628/2, Ground floor, Behind CNG Station,
M.G Road, Lado Sarai, New Delhi-110 030.

Subject: A Multi- centre, randomized, double masked, placebo controlled, parallel group, phase-III comparasion study to access efficacy and safety of Bibrocathol 2% Eye Ointment (Posiformin 2%) in the treatment of Chronic Blepharoconjunctivitis.- regarding

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Reference: Your application dated 25.04.2018 on the subject mentioned above.

Sir,
This Directorate has no objection to your conducting the subject mentioned clinical trial as per the provisions of Drugs & Cosmetics Rules under supervision of the following investigator and as per the **Protocol No: CT-POSI-1001-2016 Version No: 01, Dated 08.02.2017** submitted to this Directorate.

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| 1 | Dr. Anand Balasubramanian | Sankara Eye Hospital, Kundalhali gate, Varthur main road, Bangalore-560037 | Ethics committee Institutional review board, Sri Kanchi Kamakoti Medical trust Sankara Eye Hospital, Kundalhali gate, Varthur main road, Bangalore-560037 ECR/705/Inst/KA/2015 |
| 2 | Dr. Sanjay Verma | PARAS Hospital, C-1, Sushant Lok-1, Sctor-43, Guragaon, Haryana-122002 | PARAS Hospital, C-1, Sushant Lok-1, Sctor-43, Guragaon, Haryana-122002 ECR/249/Inst/HR/2013/RR-16 |
| 3 | Dr. Abhisek Kumar Gupta | Nirmal Hospital, Opp. MLB mediollege, gate no.3, Jhansi (U.P.),284128 | Nirmal Hospital Institutional Ethics Committee, Nirmal Hospital, Opp. MLB medical College, gate no.3, Jhansi (U.P.),284128 ECR/325/Inst/UP/2013/RR-16 |

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| 4 | Dr. Rajat Kumar Srivastava | King George Medical University, Lucknow-226003 | Institutional Ethics Committee, Research Cell, Administrative Block, KGMU, Lucknow-226003 ECR/262/Inst/UP/2013 |
| 5 | Dr. Sandeep b Patil | Vijya Ortho and Trauma Centre, Opp. BIMS, Vijaya Road, Ayodhya Nagar, Belgaum-01, Karnataka | Vijya Ortho and Trauma Centre (VOTC), Ethics Committee, Opp. BIMS, Vijaya Road, Ayodhya Nagar, Belgaum-01, Karnataka ECR/781/Inst/KA/2015 |
| 6 | Dr Smitha.K.S | KLE, Dr. Prabhakara Kore Hospital and Medical Research Centre, Nehru Nagar, belagavi-590010, Karnataka, India | Institutional Ethics Committee, KLES, Dr. Prabhakara Kore Hospital and Medical Research Centre, Nhru Nagar, belagavi-590010, Karnataka, India ECR/211/Inst/KA/2013/RR-16 |
| 7 | Dr. V. Vijaya Lakshmi | Dept. Of Ophthalmology, New Govt. Hospital (Associated by Govt. Siddhartha medical College)Ring road, Gunadala, Vijayawada-520008, A.P. India | Institutional Ethics Committee, Siddhartha medical College & Govt. General Hospital (IEC SMC & GGH), Govt. Siddhartha Medical Campus, Ring Road, Gunadala, Vijayawada-520008, Andhra Pradesh, India ECR/633/Inst/AP/2014/RR-17 |
| 8 | Dr. Nita Gurha | Head of Department & Sr. Consultant, Department of Ophthalmology, Batra Hospital, New Delhi-62 | Scientific research & Review Committee(SRERC), Department of Laboratory Medicine, Batra Hospital & Medical Research Centre 1, Tuglakabad Instiitutional Area, M.B. Road, New Delhi-110062, India ECR/295/Inst/DL/2013/RR-16 |
| 9 | Dr. V. Sreekumar | Kakatiya Research Centre, Mahatma Gandhi Memorial Hospital, Mahatma Gandhi Road, Warangal, Telangana-506007 | Kakatiya Institutional Ethics Committee, Kakatiya Medical Colleg, Rangampet Street, Warangal-506007, Telangana, India ECR/1469/Kakatiya/Inst/TG/2016 |

Kindly note that the clinical trial permission is subject to the following conditions:

- a) Clinical trial shall be conducted in compliance with the approved protocols, requirements of Schedule Y, Good Clinical Practice Guidelines issued by this Directorate and other applicable regulations.
- b) Approval of Institutional Ethics Committee duly registered with CDSCO (under Rule 122DD of Drugs & Cosmetics Rules) should be obtained and submitted to this Directorate before initiation of the study.

- c) Clinical trials shall be registered at Clinical Trials Registry of India (CTRI) before enrolling the first patient for the study.
- d) Annual status report of each clinical trial, as to whether it is ongoing, completed or terminated, shall be submitted to the Licensing Authority, and in case of termination of any clinical trial the detailed reasons for the same shall be communicated to the said Licensing Authority.
- e) Any report of serious adverse event occurring during clinical trial study to the subject, after due analysis, shall be forwarded within fourteen days of its occurrence as per Appendix XI and in compliance with the procedures prescribed in Schedule Y.
- f) In case of an injury or death during the study to the subjects, the applicant shall provide complete medical management and compensation in the case of trial related injury or death in accordance with rule 122 DAB and the procedures prescribed under Schedule Y, and the details of compensation provided in such cases shall be intimated to the Licensing Authority within thirty days of the receipt of the order of the said authority.
- g) The premises of Sponsor including their employees, subsidiaries and branches, their agents, contractors and sub-contractors and clinical trial study sites shall be open to inspection by the officers authorized by the Central Drugs Standard Control Organization, who may be accompanied by an officer of the State Drug Control Authority concerned, to verify compliance to the requirements of Schedule Y, Good Clinical Practices guidelines for conduct of clinical trial in India and other applicable regulations.
- h) The Sponsor including their employees, subsidiaries and branches, their agents, contractors and sub-contractors and clinical trial study sites and the Investigator shall allow officers authorized by the Central Drugs Standard Control Organization, who may be accompanied by an officer of the State Drug Control Authority concerned, to enter with or without prior notice, any premises of Sponsor including their employees, subsidiaries and branches, their agents, contractors and sub-contractors and clinical trial sites to inspect, search and seize any record, data, document, books, investigational drugs, etc. related to clinical trial and provide adequate replies to any queries raised by the inspecting authority in relation to the conduct of clinical trial.
- i) Clinical trial shall be conducted only at those sites which are institutes/hospitals having adequate emergency facilities and duly registered Institutional Ethics committees.
- j) The sponsor shall ensure that the number of clinical trials an investigator can undertake should be commensurate with the nature of the trial, facility available with the Investigator etc.

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- k) The details of payment/honorarium/financial support/fees paid by the Sponsor to the Investigator(s) for conducting the study shall be made available to this directorate before initiation of each of the trial sites.
- l) In addition to the requirement of obtaining written informed consent, an audio-video recording of the informed consent process in case of vulnerable subjects in clinical trial of New Chemical Entity or New Molecular Entity including procedure of providing information to the subject and his understanding on such consent, shall be maintained by the investigator for record; provided that in case of clinical trial of anti-HIV and anti-leprosy drugs, only audio recording of the informed consent process of individual subject including the procedure of providing information to the subject and his understanding on such consent shall be maintained by the investigator for record, as per Government of India, Gazette Notification vide G. S. R. no. 611(E) dated 31.07.2015.
- m) The bulk drug to be used in manufacturing of finished formulation intended to be used in the clinical trial and clinical trial batches of finished formulation shall be manufactured under GMP conditions using validated procedures and shall have ongoing stability Programme.
- n) If the clinical trial batches are different from that of the primary batches for which data have been submitted, stability reports for clinical trial batches are to be submitted as per Appendix IX of schedule Y of drugs and Cosmetics Rules for Drug substances and formulation along with Clinical Study Report.
- o) It may kindly be noted that merely granting permission to conduct clinical trials with the drug does not convey or imply that based on the clinical trial data generated with the drug, permission to market this drug in the country will automatically be granted to you.
- p) Bioequivalence with the reference product shall be established before initiation of the clinical trial.
- q) Informed consent documents (ICD) viz. Patient Information Sheet (PIS) and Informed Consent Form (ICF) complete in all respect as per the requirements specified in Appendix V of Schedule Y of the Drugs and Cosmetics Rules, 1945 must be got approved from respective Ethics Committee and submitted to CDSCO before enrolling first subject at the respective site.

Yours faithfully,



(Dr. S. Eswara Reddy)
Drugs Controller General (India)
and Licensing Authority