

6521/27.02.2015

**GOVERNMENT OF INDIA**

Directorate General of Health Services  
 Central Drugs Standard Control Organization  
 Office of Drugs Controller General (India)  
**(Global Clinical Trial Division)**  
 FDA Bhawan, Kotla Road, New Delhi-110002  
 Tel No: 01123236965, Fax: 01123236971  
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File No: CT/05/15 - DCG (I)

Date: 28/09/15

To,

M/s Axis Clinicals Limited  
 1-121/1, Miyapur, Hyderabad-500 049.

**Subject:** Permission for conducting a **clinical endpoint bioequivalence study** entitled **"A Randomized Double-Masked, Parallel Group, Placebo Controlled, Multicentric Study to Evaluate the Safety and Efficacy of Linaclotide Capsules 290 MCG in Subjects of Irritable Bowel Syndrome with Constipation [Group-1: Linaclotide Capsules MCG (Test), Group II: Linzess® Linaclotide Capsules 290 MCG (Reference) and Group III : Placebo]."** – regarding.

**Reference:** - Your letter no. ACL/SP-001/2015/013 dated 17 Feb 2015 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 investigators and as per the **Protocol No: CR039-14 version 1.0 dated 12 Jan 2015** submitted to this Directorate.

1. Dr. Prabhakar, Osmania General Hospital, Dept. of Gastroenterology, Afzalgunj, Hyderabad-500012.
2. Dr. Shravan Kumar, Gandhi Hospital, Dept. of Gastroenterology, No. 6-1-34, Bhoiguda Road, Musheerabad, Secunderabad-500003.
3. Dr. Gaurav Kumar Gupta, Dept. of Gastroenterology, Sawai Man Singh Hospital, JLM Marg, Jaipur-302004, Rajasthan.
4. Dr. Deepak N Amarapurkar, Bombay Hospital & Medical Research Centre, Room No.5, MRC, 12 New Marine lines, Mumbai- 400020.
5. Dr. Gopala Krishna Dhali, School of digestive & liver disease, IPGME & R, 244 A, J.C Bose Road, Kolkatta-700020.
6. Dr. Abhijit Chandra, Dept. of Gastroenterology, King George's Medical University, Lucknow Chowk, Lucknow-226003, UP.
7. Dr. V.K. Dixit, Dept. of Gastroenterology, Institute of Medical Sciences, Banaras Hindu University, Lanka, Varanasi-221005, UP.
8. Dr. Hemant Gupta, Samvedna Hospital, B27/88G, Ravindrapuri, Varanasi-221005, UP.
9. Dr. Ashwin Porwal Inamdar Multispeciality Hospital, Hospital building, S.No 15, Fatima Nagar, Pune-411040, MH-India.
10. Dr. Manish Bhatnagar, Sterling Hospital, Sterling Hospital Road, Memnagar, Ahmedabad-380 052, Gujarat, India.
11. Dr. Sanjay Rajput, Kaizen Hospital, 132 ft ring road, Helmet Circle, Memnagar, Ahmedabad-380052, Gujarat, India.

12. Dr. Narang Sushil G.L. SAL Hospital and Medical Institute, Opposite Doordarshan, Drive in Road, Ahmedabad-380054, Gujarat, India.
13. Dr. Kalidas Biswas, Medical College & Hospitals, Kolkatta, MCH Building, 4th floor 88, College Street, Kolkatta-700073.
14. Dr. Ravi Shankar, Yashoda Hospital, Behind Hari Hara Kala Bhavan, SP Road, Secunderabad-500003.
15. Dr. Saumin Shah, Unique Hospital- Multispeciality & Research Institute, Opp. Kiran Motor, Nr.Canal, Civil Hospital Char Rasta-Sosyo Circle Lane, Off ring Road, Surat-395002, Gujarat- India.
16. Dr. Pankaj Jain, Sterling Hospital, Opp. Inox, Course Circle (west), Vadodara, Gujarat, India.
17. Dr. Murali Krishna, Dept. of Gastroenterology, King George Hospital, Maharanipeta, Visakhapatnam-530002, AP

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

- a. **The standard of care should be defined under protocol.**
- b. **The trial centers should be evenly distributed across the country.**
- c. 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;
- d. Approval of the Ethics Committee shall be obtained before initiation of the study;
- e. Clinical trials shall be registered at Clinical trials Registry of India before enrolling the first patient for the study;
- f. 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;
- g. Any report of serious adverse event occurring during clinical trial study to the subject, after due analysis, shall be forwarded within 14 days of its occurrence as per Appendix XI and in compliance with the procedures prescribed in Schedule Y;
- h. 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;
- i. 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;

- j. 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.
- k. Clinical trial shall be conducted only at those sites which are institutes/hospitals having adequate emergency facilities and duly registered Institutional Ethics committees.
- l. 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. However, under no circumstances the number of trials should be more than three at a time.
- m. 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.
- n. An audio-video recording of the informed consent process in case of vulnerable subjects in clinical trials 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
- 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.

Yours faithfully,

(Dr. G.N. Singh)  
Drugs Controller General (India)

