

GOVERNMENT OF INDIA

19947/24/06/2016

Directorate General of Health Services
Central Drugs Standard Control Organization
(Global Clinical Trial Division)

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File No: CT/27/16-DCG (I)

Date: 15/03/17

To,
M/s. Roche Products (India) Pvt. Ltd.,
1503, 15th Floor, "The Capital", Plot No. C-70,
Behind ICICI Bank, BandraKurla Complex,
Bandra (E), Mumbai-400 051, India.

Subject: Permission for conducting "Phase III, Randomized, Double-Blind, Placebo-Controlled, Multicenter Study to Evaluate the Efficacy (Maintenance of Remission) and Safety of Etrolizumab compared with Placebo in Patients with Moderate to Severe Active Ulcerative Colitis who are naïve to TNF inhibitors".— regarding.

Reference: Your letter No. REG/2016/CTA/137 dated 21/06/2016 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 investigators mentioned in below and as per the Protocol No: GA229102, version no: 5, date: 28-aug-2015 submitted to this Directorate.

1. Dr. Shrikant Mukewar, Midas Multispecialty Hospital Pvt. Ltd. Maharashtra.
2. Dr. Nachiket Dubale, Poona Hospital and Research Center, 27, Sadashiv Peth, Pune-411030, Maharashtra.
3. Dr. B. Ramesh Kumar, Osmania Medical Hospital Hyderabad-500012, Telangana.
4. Dr. Ashok Jhajharia, SMS Medical College and Attached Hospital's J.L.N. Marg, Jaipur Rajasthan-302004.
5. Dr. Saumin Prakashbai Shah, Nirmal Hospital Pvt. Ltd., Ring Road, Surat Gujarat-395002.
6. Dr. Kalla Mukesh, S R Kalla Memorial Gastro and General Hospital, 78-79 Dhuleshwar Garden Behind HSBC Bank Sardar Patel Marg C-Scheme Jaipur-302001, Rajasthan.
7. Dr. Chetan Mehta, Shree Giriraj Multispecialty Hospital, 27-Navjyot Park Corner, 150 feet Ring Road, Rajkot-360 005, Gujarat.
8. Dr. Varadaraj Gokak, KLES Dr. Prabhakar Kore Hospital and Medical Research Centre, Nehru Nagar, Belagavi-590010, Karnataka.
9. Dr. Philip Abraham, P.D. Hinduja National Hospital and Medical Research Centre, Veer Savarkar Marg, Mahim, Mumbai-400016.
10. Dr. Tantry. B. Vishwanath, KMC HOSPITAL Dr. B.R. Ambedkar Circle Mangaluru-575001, Karnataka Satellite site Manipal Centre for Clinical Research 7th Floor, KMCHospital Attavar, Mangalore.
11. Dr. Sarat Chandra Panigrahi, Sparsh Hospital and Critical care Pvt. Ltd. A/407, Back Side of Kalyan Jewellers, Sahid Nagar, Bhubaneswar, Odisha.

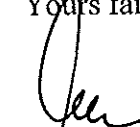
11. Dr. Sarat Chandra Panigrahi, Sparsh Hospital and Critical care Pvt. Ltd./A/407, Back Side of Kalyan Jewellers, Sahid Nagar, Bhubaneswar, Odisha.
12. Dr. Sobna Bhatia, Office Room No. 1118, MS Building, 11th Floor, Seth G.S. Medical College & KEM Hospital, Parel, Mumbai-400012.
13. Dr. Moh. Aejaz Habeeb, Centre for Liver Research & Diagnostics dept. of Gastroenterology & Hepatology, Deccan College of Medical Science, Owaisi Hospital Hyderabad-500058.
14. Dr. Nitin Pai, Grant Medical Foundation Ruby Hall Clinic, Pune-411001
15. Dr. Arijit De, Institute of Digestive and Liver Diseases, Dispur Hospital Pvt. Ltd., Ganeshguri Guwahati-781006, Assam.

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 the Ethics Committee shall be obtained before initiation of the study.
- c. Clinical trials shall be registered at Clinical trials Registry of India 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 14 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 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.
- k. 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.
- l. 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 (I)

