

GOVERNMENT OF INDIA

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

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File number: CT/17/000066

To,

M/s. Boehringer Ingelheim India Pvt. Ltd.,
1102, 11th Floor, Hallmark Business Plaza,
Guru Nanak Hospital Road, Near Guru Nanak Hospital,
Bandra (East), Mumbai-400 051.

Subject: Clinical trial titled “An open-label extension trial to assess the long term safety of Nintedanib in patients with ‘Systemic Sclerosis associated Interstitial Lung Disease’ (SSc-ILD)” – regarding.

Reference: Your online application reference no. GCT/Form44/FF/2017/4417 (GCT /68/17) dated 23/08/2017.

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: 1199.225, Version 1** dated 30 /May/2017 submitted to this Directorate.

1. Dr. Swamakar Rajesh Nathuram, Getwell Hospital & Research Institute, 20/1, Dr. Khare Marg, Dhantoli, Nagpur-440012.
2. Dr. Balamugesh Thangakunam, Dept. of Pulmonary Medicine, Christian Medical College, Ida Scudder Road, Vellore - 632004, India.
3. Dr. Zarir F. Udawadia, Pulmonary Function Lab, P.D Hinduja National Hospital & Medical Research Centre, Veer Savarkar Marg, Mahim West, Mumbai-400016, Maharashtra.
4. Dr. Mahendra Mangilal Kawedia, Jehangir Clinical Development Centre Pvt. Ltd. , Jehangir Hospital Premises, 32 Sassoon Road, Pune-411001, Maharashtra, India.
5. Dr. Jugal Kishore Kadel, Consultant Rheumatologist, Care Hospitals, Road No. 10, Banjara Hills, Hyderabad-500034, Telangana, India
6. Dr. Digamber Behara, Dept. of Pulmonary Medicine, 4th Floor, F- Block, Room No. 11, Nehru Building- PGIMER, Sector-12, Chandigarh - 160012..
7. Dr. Ujjwal Kumar Parakh, Consultant, Chest Medicine /Room No. G8 A, Ground Floor, Sir Ganga Ram Hospital, Sir Ganga Ram Hospital Marg, Rajinder Nagar, New Delhi- 110060.
8. Dr. Uma Kumar, Professor & Head, Dept. of Rheumatology, Room No 4076, 4th Floor, Teaching Block, AIIMS, New Delhi.
9. Dr. Vineeta Shobha, Professor Head, Dept. of Clinical Immunology & Rheumatology, St. John's Medical College Hospital, Koramangala, Sarjapur Road, Bangalore-560034.
10. Dr. Lokhande Rahul Maruti, Associate Professor, Dept. of TB and Chest, Department of TB and Chest, B. J. Government Medical College, and Sassoon General Hospitals, Somwar Peth, Pune - 411001, Maharashtra, India.

11. Dr. Murali Mohan B.V, Consultant Pulmonologist, Mazumdar Shaw Medical Centre (Unit of Narayan Health), #258/ A, Bommasandra Industrial Area, Anekal Taluk, Bangalore 560099, Karnataka, India.
12. Dr. Gayathri Devi H.J., Prof & Head Dept. of Pulmonary Medicine, M S Ramaiah Medical College and Hospitals , M S Ramaiah Nagar, MSRIT Post, Bangalore-560054, Karnataka, India.
13. Dr. Devarasetti Phani Kumar, Asst. Professor- Dept. of Rheumatology, 4thFloor, Millennium Block, Dept. of Rheumatology, Nizam's Institute of Medical Sciences, Panjagutta, Hyderabad, Telangana- 500082, India.
14. Dr. Ashish Kumar, Sr. Pulmonologist, Asthma Bhawan, R-3, Sector-6, Vidyadhar Nagar, Jaipur- 302039, Rajasthan.
15. Dr. Ajit Bapurao Nalawad, Consultant Rheumatologist, Inamdar Multispecialty Hospital, Hospital Building, S. No. 15, Behind KPCT Mall, Fatima Nagar, Pune, Maharashtra-411040.

Note that the clinical trial permission is subject to the following conditions:

- a. **Chest x-ray and microbiological assessments must be performed at regular intervals and or optimal diagnosis and management of TB and Pneumonia.**
- b. 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.
- c. Approval of the Ethics Committee shall be obtained before initiation of the study.
- d. Clinical trials shall be registered at Clinical trials Registry of India before enrolling the first patient for the study.
- e. 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.
- f. 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.
- g. 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.
- h. 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.
- i. 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.
- j. Clinical trial shall be conducted only at those sites which are institutes/hospitals having adequate emergency facilities and duly registered Institutional Ethics committees.

- 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. 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.
- m. 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.
- n. 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 the respective Ethics committee and submitted to CDSCO before enrolling first subject at the respective site.

Yours faithfully,

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