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**File No CT/212/12-DCG(I)** **Clinical Trial NOC No:- GCT/19/13**

53531/25/10/13

**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
Te No: 01123236965, Fax: 01123236971
E-mail: dci@nb.nic.in, cdscog@gmail.com

**File No: CT/212/12-DCG (I)** Dated: 4/12/13

To

M/s. Eli-Lilly & Company (India) Pvt. Ltd.,
Plot No. 92, Sector-32
Gurgson-122001, Haryana

**Subject: Permission for conducting a phase III clinical study entitled, "A Randomized, Double-Blind, Placebo-Controlled, Phase 3 Study Evaluating the Efficacy and Safety of Baricitinib (LY3009104) in Patients with inadequate response to conventional disease –modifying antirheumatic drug with Severely Active Rheumatoid Arthritis."** – regarding.

**Clinical Trial NOC No.** GCT/19/13

**Reference**: Your letter no. LILLY/RA/DCGI/2013/208 dated 18/10/13 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: I4V-MC-JADX (a) dated 21-08-2012** submitted to this Directorate.

1. Dr Milind Nadkar, Seth GS Medical College and KEM Hospital, Acharya Donde Marg, Parel, Mumbai – 400012.
2. Dr Jyotsana Oak, Kokilaben Dhirubhai Ambani Hospital & Medical Research Institute – Mumbai – 400053
3. Dr. Chandrasekara Srikantiah, Chanre Rheumatology & Immunology Centre for Research, No:149, 15th Main, NHCL Water Tank Road, 4th Block, 3rd Stage, Basaveshwarnagar, Bangalore – 560079
4. Dr. Vineeta Shobha, St. John's Medical College & Hospital, Sarjapur Road, Bangalore – 560034

5. Dr. Reena Sharma, SHALBY Hospital, Clinical Research Dept, 9th Floor, Opp Karnavati Club, SG Highway, Ahmedabad – 380015
6. Dr. Padmanabha Shenoy D, AMRITA Institute of Medical Sciences and Research Centre, AIMS Ponekkara Post, Kochi, Kerala – 682041
7. Dr. Alakendu Ghosh, Institute of Postgraduate Medical Education and Research, 244, AJC Bose Road, Ronald Ross Building, 4th Floor, Room Number – 7, Kolkata
8. Dr. Sarath Chandra Mouli Veeravalli, Dept. of Rheumatology, KIMS Hospital, #1-8-31/1, Ministers Road, Secunderabad-500003
9. Dr. Sarvajeet Pal, Sumana Hospital, Plot Number: 687,688, Vivekanand nagar Colony, Kukatpally, Hyderabad – 500072
10. Dr. Rajiva Gupta, Department of Rheumatology and Clinical Immunology, Medanta The Medicity, Sector 38, Gurgaon, Haryana 122 001
11. Dr. Sundeep Kumar Upadhyaya, Department of Rheumatology, Indraprastha Apollo Hospitals, Sarita Vihar, Delhi-Mathura Road, New Delhi 110076.
12. Dr. D C Sundaresh, M S Ramaiah Medical College & Hospital, New BEL Road, MSRIT Post, Bangalore - 560054
13. Dr. Surendra U kamath, Manipal Centre for Clinical Research, 7th Floor, K M C Hospital, Attavar, Mangalore – 575003
14. Dr. Shyamasish Bandhoypadhya, Apollo Gleneagles Hospitals, 58 Canal Circular Road, Kolkata 700054
15. DR. Jugal Kishore Kadel, Gurunanak CARE Hospital, 1-4-908/7/1, Bakaram, Musheerabad Main Road, Musheerabad, Hyderabad 500020, Andhra Pradesh
16. Dr. Avinash Aggarwal, Shri Nidan Hospital and Hope Fertility Centre, 27 Vidhut Nagar-A, Ajmer Road, Jaipur 302 006, Rajasthan.
17. Dr Able Lawrence, Department of Clinical Immunology, Sanjay Gandhi Post Graduate Institute, Raibarely Road, Lucknow, Uttar Pradesh
18. Dr. Siddhartha Das, Professor and Head, Department of Rheumatology, King George Medical College – Lucknow – 226003, Uttar Pradesh
19. Dr. Shailesh V Udapudi, Department of Orthopaedics, KLES Dr. Prabhakar Kore Hospital & MRC, J N Medical College, Belgaum, Karnataka – 590010

20. Dr. Piyush Parikh, AMC MET Medical College and Seth LG General Hospital, Rambag, Maninagar, Ahmedabad - 380008, Gujarat

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 ten 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. However, under no circumstances the number of trials should be more than three at a time.

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. You are also informed that audio visual recording of the informed consent process of each trial subject, including the procedure of providing information to the subject and his/her understanding on such consent is required to be done duly adhering to the principles of confidentiality. Such audio-visual recording and related documentation shall be preserved.

m. All the sponsors /investigators /institutes/Organizations and other stake holders involved in conduct of clinical trials in the country are hereby directed to adhere to the above requirement of audio-visual recording of informed consent process of trial subjects with immediate effect.

n. 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)**