

**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  
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90/02/01/13

File No: CT/143/12-DCG (I)

Dated: 20 AUG 2013

To,  
M/s. Asiatic Clinical Research Pvt. Ltd.,  
169/53, 14 Main Road, 1<sup>st</sup> Block East,  
Jayanagar, Bangalore-560011.

**Subject:** Clinical trial (Phase-III) with **ART-123 (recombinant soluble Thrombomodulin alfa -6 mg/ml in glass ampoules)** as per **Protocol No: 3-001 version 1.0 dated 23 May 2012**; - regarding.

**Clinical Trial Permission No:** GCT/02/13

**Reference:** Your letter no. CT/SEP/DCGI/2012/03 dated 31.12.2012 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 below.

1. Dr. Kattadiyil Poulouse, SUT Hospital Pattom P.O, Post Box-1052 Trivandrum, Kerala 695004
2. Dr. Mukesh Kumar Sarna, Medanta- Monilek Hospital and Research Centre, Sec-4, Jawahar Nagar Jaipur Rajasthan 302004
3. Dr. Mradul kumar daga, Maulana Azad Medical College and Associated, L N Hospital, 129, B. L. Taneja Block, Bahadur Shah Zafar Marg New Delhi 110002
4. Dr. Parasuram Krishnamoorthi Meenakshi Mission Hospital and Research Centre, Lake Area, Melur Road Madurai Tamilnadu 625107
5. Dr. Sameer Jog Deenanath Mangeshkar Hospital and Research Centre, Erandawane Pune Maharashtra
6. Dr. Pradeep Micheal D'costa King Edward Memorial Hospital 489, Rasta Peth, Moodliar Road Pune Maharashtra 411011.
7. Dr. Sarat Kumar Behera Hi-tech Medical College & Hospital Pandara, Rasulgarh Bhubaneshwar Odisha 751010.

This 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 ;( Please refer [www.ctri@nic.in](http://www.ctri@nic.in)
- d. Annual status report of each clinical trial, as to whether it is ongoing, completed or terminated, shall be submitted to the Licensing Authority under Rule 21(b)[i.e. DCG(I)] 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. Before initiation of the study, you are required to obtain approval of Ethics Committee which is duly registered under the Provision of Drugs and Cosmetics Rules 1945.
- 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. Upper age limit of the subjects should be 65 years.

Yours faithfully,

  
(Dr. G.N. Singh)

**Drugs Controller General (India)**

Copy forwarded for Information

1. Drugs controller (Karnataka), State Drug control Department, PB No. 5377, Palace Road, Bangalore-560001.
2. Asstt. Drugs Controller (India), Palace Road, Bangalore-560001.