

IND/CT04/FF/2019/16502

F. No. IND/CT/19/000042

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Government of India
Directorate General of Health Services
Central Drugs Standard Control Organization
(New Drugs Division)

FDA Bhawan, Kotla Road,
New Delhi-110002

Dated: **11 4 NOV 2019**

To
M/s Aurigene Discovery Technologies Limited,
39-40, KIADB Industrial Area,
Phase II Electronic City,
Hosur Road,
Bengaluru - 560 100, India

Subject:- Permission for conducting clinical study entitled, "A Phase II, Multicenter, Double-blind, Double-dummy, Placebo controlled, Randomized, Study to Evaluate the Efficacy and Safety of two doses of AUR101 in patients with Moderate-to-Severe Psoriasis (INDUS-2)" - regarding.

Sir,

With reference to your Application No. IND/CT/19/000042, please find enclosed herewith the permission in Form CT-06, No. **CT/ND/99/2019** to conduct the subject mentioned clinical trial under the provisions of **New Drugs and Clinical Trial Rules, 2019.**

This permission is subject to the conditions, as mentioned below.

Yours faithfully



(Dr. V. G. Somani)
Drugs Controller General (India)
Central Licensing Authority

Condition of permission

- (i) Clinical trial at each site shall be initiated after approval of the clinical trial protocol and other related documents by the Ethics Committee of that site, registered with the Central Licencing Authority under Rule 8;
- (ii) Where a clinical trial site does not have its own Ethics Committee, clinical trial at that site may be initiated after obtaining approval of the protocol from the Ethics Committee of another trial site; or an independent Ethics Committee for clinical trial constituted in accordance with the provisions of Rule 7:

Provided that the approving Ethics Committee for clinical trial shall in such case be responsible for the study at the trial site or the centre, as the case may be:

Provided further that the approving Ethics Committee and the clinical trial site or the bioavailability and bioequivalence centre, as the case may be, shall be located within the same city or within a radius of 50 kms of the clinical trial site;

- (iii) In case an ethics committee of a clinical trial site rejects the approval of the protocol, the details of the same shall be submitted to the Central Licensing Authority prior to seeking approval of another Ethics Committee for the protocol for conduct of the clinical trial at the same site;
- (iv) The Central Licencing Authority shall be informed about the approval granted by the Ethics Committee within a period of fifteen working days of the grant of such approval;
- (v) Clinical trial shall be registered with the Clinical Trial Registry of India maintained by the Indian Council of Medical Research before enrolling the first subject for the trial;
- (vi) Clinical trial shall be conducted in accordance with the approved clinical trial protocol and other related documents and as per requirements of Good Clinical Practices Guidelines and the provisions of these rules;
- (vii) Status of enrolment of the trial subjects shall be submitted to the Central Licencing Authority on quarterly basis or as appropriate as per the duration of treatment in accordance with the approved clinical trial protocol, whichever is earlier;
- (viii) Six monthly status report of each clinical trial, as to whether it is ongoing, completed or terminated, shall be submitted to the Central Licencing Authority;
- (ix) In case of termination of any clinical trial the detailed reasons for such termination shall be communicated to the Central Licencing Authority within thirty working days of such termination;
- (x) Any report of serious adverse event occurring during clinical trial to a subject of clinical trial, shall, after due analysis, be forwarded to the Central Licencing Authority, the chairperson of the Ethics Committee and the institute where the trial has been conducted within fourteen days of its occurrence as per Table 5 of the Third Schedule and in compliance with the procedures as specified in Chapter VI of the New Drugs and Clinical Trials Rules, 2019;
- (xi) In case of injury during clinical trial to the subject of such trial, complete medical management and compensation shall be provided in accordance with the Chapter VI of the said Rules and details of compensation provided in such cases shall be intimated to the Central Licencing Authority within thirty working days of the receipt of order issued by Central Licencing Authority in accordance with the provisions of the said Chapter;
- (xii) In case of clinical trial related death or permanent disability of any subject of such trial during the trial, compensation shall be provided in accordance with the Chapter VI and details of compensation provided in such cases shall be intimated to the Central Licencing Authority within thirty working days of

receipt of the order issued by the Central Licencing Authority in accordance with the provisions of the said Chapter;

- (xiii) The premises of the sponsor including his representatives and clinical trial sites, shall be open for inspection by officers of the Central Licencing Authority who may be accompanied by officers of the State Licencing Authority or outside experts as authorised by the Central Licencing Authority, to verify compliance of the requirements of these rules and Good Clinical Practices Guidelines, to inspect, search and seize any record, result, document, investigational product, related to clinical trial and furnish reply to query raised by the said officer in relation to clinical trial;
- (xiv) Where the New Drug or Investigational New Drug is found to be useful in clinical development, the sponsor shall submit an application to the Central Licencing Authority for permission to import or manufacture for sale or for distribution of new drug in India, in accordance with Chapter X of these rules, unless otherwise justified;
- (xv) The Laboratory owned by any person or a company or any other legal entity and utilised by that person to whom permission for clinical trial has been granted used for research and development, shall be deemed to be registered with the Central Licensing Authority and may be used for test or analysis of any drug for and on behalf of Central Licensing Authority;
- (xvi) The Central Licencing Authority may, if considered necessary, impose any other condition in writing with justification, in respect of specific clinical trials, regarding the objective, design, subject population, subject eligibility, assessment, conduct and treatment of such specific clinical trial;
- (xvii) The sponsor and the investigator shall maintain the data integrity of the data generated during clinical trial.
- (xviii) Informed Consent Documents (ICD) viz. Patient Information Sheet (PIS) and Informed Consent Form (ICF) complete in all respect & must be got approved from the respective Ethics committee and submitted to CDSCO before enrolling first subject at the respective site.

FORM CT-06

(See rules 22, 25, 26, 29 and 30)

**PERMISSION TO CONDUCT CLINICAL TRIAL OF NEW DRUG OR INVESTIGATIONAL
NEW DRUG**

The Central Licensing Authority hereby permits M/s Aurigene Discovery Technologies Limited, 39-40, KIADB Industrial Area, Phase II Electronic City, Hosur Road, Bengaluru - 560 100, India to conduct clinical trial of the investigational new drug as per **Protocol No. AUR101-201, Version No. 1.0, Dated 07.08.2019** in the below mentioned clinical trials sites.

2. Details of new drug or investigational new drug and clinical trial site:-

Names of the new drug or investigational new drug:	AUR101		
Therapeutic class:	Anti-inflammatory		
Dosage form:	Tablets		
Composition:	1. AUR101 Tablets 100mg Each tablet contains: AUR101 100mg. 2. AUR101 Tablets 300mg Each tablet contains: AUR101 300mg.		
Indications:	Psoriasis		
Details of clinical trial sites-			
Sr. No.	Name of Principal Investigator & Trial Sites	Ethics Committee Name/ Registration Number	
01	Dr. Shyamal Balki, Shree Hospital and Critical Care Centre, 799, OM Nagar, Opp. Tajshree Building Sakkardara Sq. Nagpur-440009, Maharashtra, India	Shree Hospital Ethics Committee, Shree Hospital Unit, Plot No. 786 A, 3rd Floor, Behind Shree Hospital & Critical Care Centre. ECR/553/Inst/MH/2014/RR-17	
02	Dr. M. Kanaka Prasad Rao, Govt. Medical College & Govt. General Hospital (Old RIMSGGH), Srikakulam-532001, Andhra Pradesh, India	Institutional Ethics Committee, Rajiv Gandhi Institute of Medical Sciences & RIMS Government General Hospital, Srikakulam- 532001, Andhra Pradesh, India. ECR/492/Inst/AP/2013/RR-16	
03	Dr. Abanti Saha, Medical College Kolkata, 88 College Street, College Square, Kolkata- 700073, West Bengal, India.	Institutional Ethics Committee for Human Research, Medical College Kolkata, 88, College Street, College Square, Kolkata- 700073, West Bengal, India ECR/287/Inst/WB/2013/RR-16	


04	Dr. Abhishek De, Calcutta National Medical College, 32, Gorachand Road, Beniapukur, Kolkata-700014, West Bengal, India.	Ethics Committee, Calcutta National Medical College, 32, Gorachand Road, Kolkata-700014, West Bengal, India. ECR/771/Inst/WB/2015
05	Dr. Sunil Kumar, Dayanand Medical College and Hospital, Tagore Nagar, Civil Lines, Ludhiana-141001, Punjab, India.	Drug Trial Ethics Committee, Basement, Hero DMC Heart Institute, Opp. Library, Dayanand Medical College and Hospital, Tagore Nagar, Civil Lines, Ludhiana-141001, Punjab, India. ECR/101/Inst/PB/2013/RR-16
06	Dr. Sushil Yashwant Pande, NKP Salve Institute of Medical Sciences and Lata Mangeskar Hospital, Dighdoh Hills, Hingna, Nagpur-440019, Maharashtra, India.	Institutional Ethics Committee, NKP Salve Institute of Medical Sciences, Dighdoh Hills, Hingna, Nagpur-440019, Maharashtra, India. ECR/88/Inst/MH/2013/RR-19
07	Dr. Nandini A S, Kempegowda Institute of Medical Sciences, Department of dermatology, "B" Block, OPD Building, K. R. Road, V. V. Puram, Bangalore – 560004.	KIMS Institutional Ethics Committee, Banashankari II Stage, Bangalore-560070. ECR/486/Inst/KA/2013/RR-16
08	Dr. Savitha A S, Sapthagiri Hospital, Ground floor, dermatology OPD, #15, Chikkasandra, Hesaraghatta, Main Road, Bangalore –560090.	Institutional Ethics Committee, Sapthagiri Institute of Medical Science & Research Center, Chikkasandra, Hesaraghatta, Bangalore -560090. ECR/583/Inst/KA/2014/RR-17
09	Dr. Bela J Shah, B.J Medical College & Civil Hospital, Department of Dermatology, Wing No.3, Room No. 139, skin OPD 1st Floor, OPD Building, Asarwa, Ahmedabad – 380016.	Institutional Ethics Committee, BJ Medical College & Civil Hospital, Ahmedabad - 380016. ECR/72/Inst/GJ/2013/RR-19
10	Dr. Sunil Dogra, Postgraduate Institute of Medical Education & Research, Department of Dermatology, Venereology & Leprology Postgraduate Institute of Medical Education & Research (PGIMER) Chandigarh 160 012.	Institutional Ethics Committee, Postgraduate Institute of Medical Education & Research, Room No. 606, 6th Floor, PN Chuttani Block, Chandigarh – 160012. ECR/25/Inst/CH/2013/RR-16

11	Dr. Suneel Vartak, Sujata Birla Hospital & Medical Research Center, Department of dermatology, OPD No. 6, Ground Floor, Opposite to Bytco College, Nashik road, Nashik – 422101.	Yash Society Ethics Committee, Sujata Birla Hospital, Nasik - 422101. ECR/98/Inst/MH/2013/RR-19
12	Dr. Mahendra Kura, Grant Medical College and JJ Hospital, OPD-42, 2nd Floor, OPD building, JJ Hospital, Byculla, Mumbai-400008.	Institutional Ethics Committee, Grant Medical College & Sir J J Hospital Byculla, Mumbai - 400008. ECR/382/Inst/MH/2013/RR-16

3. This permission is subject to the conditions prescribed in part A of Chapter V of the New Drugs and Clinical Trials Rules, 2019 under the Drugs and Cosmetics Act, 1940.

New Delhi

Date: 14 NOV 2020


 (Dr. V. G. Somani)
 Central Licensing Authority
 Drugs Controller General (India)
 Dte. General of Health Services
 Ministry of Health and Family Welfare
 FDA Bhawan, Kotla Road, I.T.O.
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