

Diary No: 27139
Dated: 04.08.2021

F. No. ND/MA/21/000122
Government of India
Directorate General of Health Services
Central Drugs Standard Control Organization
(New Drugs Division)

Tele No.011-23236965
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FDA Bhawan, Kotla Road,
New Delhi-110002
Dated:

To

17 SEP 2021

M/s Exemed Pharmaceuticals, 133/1 & 133/2
GIDC Selvas Road, Vapi
Gujarat (India) - 396195

Subject: Grant of Permission to conduct Phase III Clinical Trial entitled "A Randomized, Double Blind, Parallel Group, Placebo Controlled, Comparative, Multicenter and Phase III Clinical Study to Evaluate the Efficacy and Safety of Bempedoic Acid Tablets 180 mg Added to Background Lipid-Modifying Therapy in Patients with Elevated LDL-C who are Statin Intolerant." - regarding.

Sir,

With reference to your application no. ND/MA/21/000122 dated 04.08.2021, please find enclosed herewith the permission in Form CT-06, No. **CT/ND/154/2021** to conduct the subject mentioned clinical trial under the provisions of **New Drugs and Clinical Trial Rules, 2019** granted based on evaluation in consultation with Subject Expert Committee (Cardiovascular and Renal).

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

Yours faithfully



(Dr. V. G. Somani)
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 authorized 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 utilized 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.
 - (xix) It may kindly be noted that merely granting permission to conduct clinical trial with the drug doesn't convey or imply that based on the clinical data generated with the drug, permission to market this drug will automatically be granted to you.

FORM CT-06

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

PERMISSION TO CONDUCT CLINICAL TRIAL OF NEW DRUG OR INVESTIGATIONAL NEW DRUGNumber of the permission and date of issue **CT/ND/154/2021** dated **17 SEP 2021**

1. The Central Licensing Authority hereby permits M/s Exemed Pharmaceuticals, 133/1 & 133/2, GIDC Selvas Road, Vapi, Gujarat (India) - 396195 to conduct clinical trial of the new drug as per **EP/BEM/CT/20/003, Version No. 01 & Dated Sep 01, 2021** in the below mentioned clinical trials sites.
2. Details of new drug or investigational new drug and clinical trial sites: -

Names of the new drug or investigational new drug:	Bempedoic Acid Tablets
Therapeutic class:	An Adenosine triphosphate citrate lyase (ACL) Inhibitor
Dosage form:	Film Coated Tablets
Composition:	Each Tablet Contains Bempedoic Acid.....180 mg Excipients.....q.s.
Indications	Indicated as an adjunct to diet and maximally tolerated statin therapy for the treatment of adults with heterozygous familial hypercholesterolemia

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: **17 SEP 2021**

(Dr. V. G. Somani)
Central Licensing Authority
Stamp

Dr. V. G. SOMANI
Drugs Controller General (India)
Dte. General of Health Services
Ministry of Health and Family Welfare
FDA Bhawan, Kotla Road, I.T.O.
New Delhi-110002

Details of clinical trial site			
SI No.	Investigator Name	Address of Site	Name of Ethics Committee and address with Ethics Committee registration No
01	Dr. Sanjiv Maheshwari	Department of Medicine, Jawahar Lal Nehru (J.L.N) Medical College, Kala Bagh, Ajmer-305001,	Institutional Ethics Committee, Jawahar Lal Nehru Medical College, Kala Bagh, Ajmer-305001,

		Rajasthan.	Rajasthan. ECR/1156/Inst/RJ/2018
02	Dr. Sanjay Sharma	Maharaja Agrasen Superspeciality Hospital, Central Spine, Agrasen Aspatal Marg, Sector 7, Vidyadhar Nagar, Jaipur-302039, Rajasthan.	Institutional Ethics Committee Maharaja Agrasen Superspeciality Hospital, Central Spine, Agrasen Aspatal Marg, Sector 7, Vidyadhar Nagar, Jaipur-302039, Rajasthan. ECR/1222/Inst/RJ/2019
03	Dr. Raja Bhattacharya	Medical College and Hospital, Kolkata, MCH Building, 4th Floor, 88 College Street, Kolkata-700073, West Bengal	Institutional Ethics Committee for Human Research Medical College, Kolkata, 88, College Street, Kolkata-700073, West Bengal. ECR/287/Inst/WB/2013/RR-19
04	Dr. Hrishikesh Bora	Down Town Hospital, Dispur, G.S. Road, Guwahati-781006, Assam.	Ethics Committee, Down Town Hospital, 6th Floor, 1st Building, G.S. Road, Dispur, Guwahati-781006, Assam. ECR/549/Inst/AS/2014/RR-20
05	Dr. Vijaykumar Bhagwan Barge	CROM Clinical Research & Medical Tourism Pvt. Ltd., CROM Clinical Research & Medical Tourism Pvt. Ltd. at Rajarshee Chhatrapati Shahu, Maharaj Govt. Medical College and Chhatrapati Pramila Raje General Hospital, Dasara Chowk, Town Hall, Bhausingji Road, Kolhapur-416002, Maharashtra.	Rajarshee Chhatrapati Shahu Maharaj Govt. Medical College and Chhatrapati Pramila Raje General Hospital, Kolhapur Institutional Ethics Committee 2 (RCSMGMCI EC2), Dasara Chowk, Town Hall, Bhausingji Road, Kolhapur-416002, Maharashtra. ECR/1052/Inst/MH/2018/RR-21
06	Dr. Vijaykumar Shivajirao Patil	Prakash Institute of Medical Sciences & Research, Urun-Islampur (PIMS&R), Tyzer Clinical Research Pvt. Ltd. at Islampur-Sangali Road, Islampur, Tal-Walwa, Dist-Sangali, Maharashtra-415409.	Prakash Medical College Institutional Ethics Committee, Prakash Institute of Medical Sciences & Research, Urun-Islampur (PIMS&R), Islampur-Sangali Road, Islampur, Tal-Walwa, Dist-Sangali, Maharashtra-415409. ECR/1052/Inst/MH/2018/RR-21
07	Dr. D. Anil Kumar	Department of General Medicine, Gandhi Medical College / Hospital, Musheerabad, Secunderabad-500003, Telangana.	Institutional Ethics Committee, Gandhi Medical College/Gandhi Hospital, Musheerabad, Secunderabad-500003, Telangana. ECR/180/Inst/AP/2013/RR-19

08	Dr. Vikas Reddy Maddali	Osmania Medical College & General Hospital, Afzalgunj, Hyderabad, Telangana-500012.	Osmania Medical College, Koti, Hyderabad-500095, Telangana State. ECR/300/Inst/AP/2013/RR-19
09	Dr. A. Gopal Rao	Government Medical College & Government General Hospital (Old RIMSGGH), Srikakulam- 532001, Andhra Pradesh.	Institutional Ethics Committee, Rajiv Gandhi Institute of Medical Sciences & RIMS Government General Hospital, Srikakulam-532001, Andhra Pradesh. ECR/492/Inst/AP/2013/RR-20

