Dairy No.:
SND/Form44/FF/15416

**F. No. SND/CT/19/000030**
**Government of India**
**Directorate General of Health Services**
**Central Drugs Standard Control Organisation**
**(Subsequent New Drugs Division)**

Tele No.011-23236965
Fax.No.011-23236973

FDA Bhawan, Kotla Road,
New Delhi-110002
Dated:

To
M/s. Dr. Reddys Laboratories Limited,
8-2-337, Road No.3, Banjara Hills,
Telangana (India), 500034.

**Subject: "Grant of permission to undertake Phase II clinical trial of New Drug - Zileuton Ointment 1.25% and 5.0%"-regarding.**

**CT NOC No.: CT/SND/104/2019**

Sir,

With reference to your Application No. SND/Form44/FF/2019/15416, please find enclosed herewith the permission in Form CT-06, CT NOC No. **CT/SND/104/2019** 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,**

V G SOMANI

Digitally signed by V G SOMANI
DN: c=IN, o=CENTRAL DRUGS STANDARD CONTROL ORGANIZATION, 2.5.4.20=111d9c06dbcb941e6f5d99ed4cea2e8412a222ecae4ab1512cbee85202ff8f, ou=DGHS,CID - 6445798, postalCode=110002, st=Delhi, cn=V G SOMANI
Date: 2019.11.08 19:11:55 +05'30'

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

## Conditions 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 electronically in the SUGAM portal;

(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 (lCF) 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~~**

**CT NOC NO.: CT/SND/104/2019**

The Central Licensing Authority hereby permits **M/s Dr. Reddys Laboratories Limited, 8-2-337, Road No.3, Banjara Hills, Telangana (India) – 500034** to conduct clinical trial of the new drug ~~or investigational new drug~~ as per **Protocol No.DFD-25-DC-001, Version: 2.0, Dated 04/Oct/2019** in the below mentioned clinical trial sites.

2. Details of new drug or investigational new drug:

| | |
|---|---|
| **Names of the new drug:** | Zileuton Ointment |
| **Therapeutic class:** | Anti-inflammatory |
| **Dosage form:** | Ointment |
| **Composition:** | 1. Zileuton Ointment 1.25%w/w<br>Each Gram of Ointment Contains<br>Zileuton …….. 12.5 mg<br>Ointment Base ……. q.s.<br><br>2. Zileuton Ointment 5.0%w/w<br>Each Gram of Ointment Contains<br>Zileuton …….. 50.0 mg<br>Ointment Base ……. q.s. |
| **Indications:** | Indicated for topical treatment of atopic dermatitis |

**Details of clinical trial sites**

| Sr. No. | Name of Principal Investigator & Trial sites | Ethics Committee Name/Registration Number |
|---|---|---|
| 1 | Dr.G. Purnima<br>New Govt. General Hospital (Associated by Govt. Siddhartha Medical College) Govt. Siddhartha Medical College Campus, Ring Road, Gunadala, Vijayawada - 520008,Andhra Pradesh, India | Institutional Ethics Committee<br>Siddhartha Medical College & Govt. General Hospital (IEC SMC & GGH)<br>Govt. Siddhartha Medical College Campus, Ring Road, Gunadala, Vijayawada – 520008<br><br>**ECR/633/INST/AP/2014/RR-17** |
| 2 | Dr. Davinder Prasad<br>Post Graduate Institute of Medical Education and Research, Department of Dermatology, Nehru Hospital Sector, Chandigarh - 160012 | Institutional Ethics Committee<br>Post Graduate Institute of Medical Education and Research, Room no. 6006, 6th floor, P N Chuttani block, Chandigarh - 160012, India.<br><br>**ECR/25/INST/CH/2013/RR-16** |
| 3 | Dr. Praneeth Kumar<br>Gleneagles Global Hospitals | Institutional Ethics Committee<br>Gleneagles Global Hospitals |

| | | |
|---|---|---|
| | 6-1-1040/1 to 4, Lakdikapool, Hyderabad, Telangana-500004, India | 6-1-1040/1 to 4, Lakdikapool, Hyderabad, Telangana-500004, India<br><br>**ECR/158/INST/AP/2013/RR-16** |
| 4 | Dr.Amit Madan<br>Ajanta Hospitals & IVF Centre<br>765, ABC Complex, Kanpur Road, Alambagh, Lucknow, Uttar Pradesh 226005 | Institutional Ethics Committee<br>Ajanta Hospitals & IVF Centre765, ABC Complex, Kanpur Road, Lucknow 226005<br><br>**ECR/611/Inst/UP/2014/RR-17** |
| 5 | Dr. Ramesh Bhat<br>Father Muller Medical College Hospital(A Unit of Father Muller Charitable Institutions), Kankanady, Mangalore, Karnataka-575002 | Father Muller Medical College'<br>Father Muller Charitable Institutions<br>Kankanady, Mangalore, Karnataka-575002<br><br>**ECR/540/Inst/KA/2014/RR-17** |
| 6 | Dr. Rajeev Aggarwal<br>M. V. Hospital and Research Centre, 314/30, Mirza mandi, Chowk, Lucknow, Uttar Pradesh 266024 | Institutional Ethics Committee<br>1st Floor of M.V. Hospital & Research Centre314/30, Mirza mandi, Chowk, Lucknow, Uttar Pradesh 266024<br><br>**ECR/13/Inst/UP/2013/RR-16** |
| 7 | Dr. S. C. Bharija<br>Sir Gangaram Hospital<br>Old Rajinder nagar, Rajinder Nagar, New Delhi, Delhi 110060 | Sir Gangaram Hospital, Ethics Committee,Room No. 1496,IV Floor,Old Rajinder nagar, Rajinder Nagar, New Delhi, Delhi 110060<br><br>**ECR/20/Inst/DL/2013/RR-16** |
| 8 | Dr. Anjeeta Dhawan<br>Maharaja Agrasen Hospital<br>West Punjabi Bagh, Delhi, 110026 | Institutional Ethics Committee<br>Room 614,6$^{th}$ Floor, Maharaja Agrasen Hospital, West Punjabi Bagh, NewDelhi-110026<br><br>**ECR/745/Inst/DL/2015/RR-18** |
| 9 | Dr. Parag Kalyani<br>Shree Hospital Siddharth Mansion, Nagar Road, Pune-411006, Maharashtra, India | Ethics Committee Shree Hospital,<br>Shree Hospital, 5$^{th}$floor, Siddharth Mansion, Nagar Road, Pune-411006, Maharashtra, India<br><br>**EC/RENEW/INST/2019/3959** |
| 10 | Dr. R Chavan<br>B. J. Govt. Medical College & Sassoon General Hospital,<br>Pune station road, Pune-411001 | Institutional Ethics Committee of B. J. Govt. Medical College and Sassoon General Hospital<br>Dept. of Pharmacology, B. J. Govt. Medical College, Sassoon road, Pune-411001, Maharashtra.<br><br>**ECR/280/Inst/Maha/2013/RR-16** |

| | | |
|---|---|---|
| 11 | Dr. Suresh Kumar Jain<br>Government Medical College<br>Rangbari Rd, Kota, Rajasthan 324010 | IEC, Government Medical College<br>Rangbari Road, Kota, Rajasthan 324010<br><br>**EC/NEW/INST/2017/1570** |
| 12 | Dr. Snehal Lunge<br>KLE Prabhakar Kore Hospital & MRC, Nehru nagar, Belgavi - 590010, Karnataka, India | Institutional Ethics Committee, KAHER, JNMC Campus Nehru nagar Belagavi-590010 Karnataka India<br><br>**ECR/211/Inst/KA/2013/RR-2016** |
| 13 | Dr. Savitha A.S.<br>Sapthagiri Institute of Medical Science & Research Centre<br>15, Hesarghatta Main Rd, Navy Layout, Chikkasandra, Chikkabanavara, Bengaluru, Karnataka 560090 | Saptagiri Institute of Medical Sciences & Research Centre Institutional ethics committee, 15, ChikkasandraHesaraghatta main road, Bangalore-560090<br><br>**ECR/538/Inst/KA/2014/RR-17** |
| 14 | Dr.R.D. Mehta<br>S.P. Medical College,Bikaner-334001Rajasthan | Institutional Ethics Committee<br>S.P Medical College, Bikaner-334001 Rajasthan<br><br>**ECR/27/SP/Inst/Raj/2013/ RR-16** |
| 15 | Dr. Vipul Gupta<br>KRM Hospital And Research Centre, 3/92-93 Vijiayant Khand Gomati Nagar, Lucknow-226010,UP, India | KRM Hospital Ethics Committee<br>3/92-93 Vijiayant Khand<br>Gomati Nagar, Lucknow-226010 UP, India<br><br>**ECR/848/Inst/UP/2016** |

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.

V G SOMANI

Digitally signed by V G SOMANI
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Date: 2019.11.08 19:11:33 +05'30'

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

**New Delhi**
**Date:**